

LONDON  
SCHOOL of  
HYGIENE  
& TROPICAL  
MEDICINE



LSHTM Research Online

Cross, JH; (2020) Iron and Infection: Neonatal Iron Transition. PhD (research paper style) thesis, London School of Hygiene & Tropical Medicine. DOI: <https://doi.org/10.17037/PUBS.04657217>

Downloaded from: <https://researchonline.lshtm.ac.uk/id/eprint/4657217/>

DOI: <https://doi.org/10.17037/PUBS.04657217>

**Usage Guidelines:**

Please refer to usage guidelines at <https://researchonline.lshtm.ac.uk/policies.html> or alternatively contact [researchonline@lshtm.ac.uk](mailto:researchonline@lshtm.ac.uk).

Available under license. To note, 3rd party material is not necessarily covered under this license: <http://creativecommons.org/licenses/by-nc-nd/3.0/>

<https://researchonline.lshtm.ac.uk>

LONDON  
SCHOOL of  
HYGIENE  
& TROPICAL  
MEDICINE



LSHTM Research Online

Cross, JH; (2020) Iron and Infection: Neonatal Iron Transition. PhD (research paper style) thesis, London School of Hygiene & Tropical Medicine. DOI: <https://doi.org/10.17037/PUBS.04657217>

Downloaded from: <http://researchonline.lshtm.ac.uk/id/eprint/4657217/>

DOI: <https://doi.org/10.17037/PUBS.04657217>

**Usage Guidelines:**

Please refer to usage guidelines at <https://researchonline.lshtm.ac.uk/policies.html> or alternatively contact [researchonline@lshtm.ac.uk](mailto:researchonline@lshtm.ac.uk).

Available under license: <http://creativecommons.org/licenses/by-nc-nd/2.5/>

<https://researchonline.lshtm.ac.uk>

**LONDON  
SCHOOL of  
HYGIENE  
& TROPICAL  
MEDICINE**



**Iron and Infection: Neonatal Iron Transition**

**James Henry Cross**

**Thesis submitted in accordance with the requirements for the  
degree of**

**Doctor of Philosophy**

**Of the**

**University of London**

**June 2020**

**Department of Population Health**

**Faculty of Epidemiology & Population Health**

**LONDON SCHOOL OF HYGIENE AND TROPICAL MEDICINE**

**Funded By:**

**Bill and Melinda Gates Foundation**

**Research Group Affiliations:**

**MRC International Nutrition Group &  
MRC Unit The Gambia at LSHTM**

## **STATEMENT OF OWN WORK**

I, James H. Cross, confirm the work presented in this thesis is my own. Where information has been derived from other sources, I confirm that this has been indicated in the thesis.



James H. Cross

13<sup>th</sup> December 2019



## ACKNOWLEDGEMENTS

I am incredibly grateful for the four years I have spent being able to conduct research at MRC Unit The Gambia at LSHTM.

My supervisor, Carla Cerami, has taught me a vast number of lessons about the world of work over the last six years. The care, compassion, and energy you have given me over half a decade have inspired me to build a career in global health research, with an emphasis on giving back to society and focusing on the people that need it most. Your scientific and technical advice has led me to try and ask the correct questions and work in a manner that gets things done. These are attitudes, which I hope to continue in my future career. I am inspired by your courage and spirit toward pushing a 'can do' attitude. Your constant availability at any day or time, your sense of humour and friendship have been invaluable when times became difficult. Thank you for all you have given me, and I hope you are proud of what we have both achieved working together.

To our theme leader and my associate supervisor, Andrew Prentice, you have overseen my work for my entire time at MRCG. Your caring comments and advice on aspects that relate to work, my career, research and life have helped me every step of the way. Thank you for accepting me into this wonderful family-like research group and supporting me in my transition from an undergraduate teenager to an individual that hopes to continue a career in global health research. Your leadership and mentorship to me and others has been inspiring.

This project required the hard work of a energetic multidisciplinary team to succeed. Many thanks to the whole of Neolnnate Study team, especially to Ousman Jarjou, all the research nurses, the Kanifing General Hospital laboratory staff, the transport team, Bubacarr JB

Touray, Sheriffo Jarjou, Nuradin Ibrahim Mohammed and Bakary Sonko. Thank you for the endless hours of high-quality work you have completed for this study.

Malcolm Velasco and Santiago Rayment Gomez, thank you for coming to The Gambia and becoming part of our team. Your hard work and everyday support were vital to the success of this study, my thesis and my sanity!

Thank you to all my collaborators during this study for your support and guidance. Thank you especially to Hal Drakesmith and Simon Andrews for being on my advisory committee. Additionally, many thanks to Andrew Armitage for providing me with excellent advice on developing multiple aspects of my review into neonatal iron metabolism.

My thanks to both of my upgrading examiners, Martin Holland and Sant-Rayn Pasricha for providing me with detailed feedback and comments, which helped shape the research project and my subsequent thesis. I would also like to thank Philip James for his career and PhD advice over the last four years.

Sarah Prentice laid the foundations for this PhD with the original pilot research study, HYPO-G. Thanks to you and the HYPO-G team, for giving me such a strong foundation to work from. To Tony Fulford and Richard Bradbury, thank you for replying to my original email back in 2013, allowing me to take a 'Year in Industry' at MRC Keneba. These early steps have led to a complete change in my career path and what I thought I could achieve in the future. Additionally, my great thanks go to Sophie Jatta and Honorary Jammeh, for looking after me on a day-to-day basis like two additional mothers, over the last four years. You have kept me moving forward and worked tirelessly to help me achieve this goal. God bless you and your families.

To my friends in The Gambia, thank you for the long nights in Mamas Restaurant over a few Julbrews discussing politics, medical research, history, wildlife, current world affairs and everything else in between. In particular, to my close and loyal Texan friend Noah, thank you for listening to my problems over the years, ever ready for long technical discussions about random research papers. Thanks also for providing me with countless laughs along the way.

Most importantly of all, my heartfelt thanks to my Mum, Dad, David, Binta, Alieu, Emmanuel, and Michael and my partner, Tianna. You have supported me through all of the toughest times of this PhD. You have celebrated all the small successes with me along the way. Without you, I would not be where I am today, and I would not have achieved this lifelong goal, that I had previously thought was out of my reach. Thank you for inspiring me, advising me, and loving me. I am incredibly grateful for the all the time, energy and love, you have given me through the years. I hope you all share the happiness of this achievement.

I want to conclude with a special mention of appreciation to my grandmother, Ann. Nan, your strength, passion and determination, along with your honest love of people, life and common sense have been guiding lights for me during this task. Our whole family would not be here without you, you're a role model for me, and I hope that you can rest and be content with what I have currently achieved, and my plans for the future.

**Note:** Thanks to Dr Noémi Roy and Dr Gladys Latunde-Dada for their guidance and advice during the examination process, along with making my viva as enjoyable and thought-provoking as possible.

## ABSTRACT

Neonates, particularly those born preterm (PTB) and low birthweight (LBW), are especially susceptible to bacterial infections that cause an estimated 225,000 deaths annually. Iron is a vital substrate for the most common organisms causing septicaemia. Full-term babies elicit an immediate post-natal hypoferraemia assumed to have evolved as an innate defence. This thesis aimed to test whether preterm and low birthweight newborns are capable of a similar response.

A longitudinal observational study was conducted in 430 hospital-delivered Gambian babies. Demographic, anthropometric and haematological data were collected from 152 babies who were either PTB (between  $\geq 32$ - $<37$  weeks gestational age) and/or LBW ( $<2500$ g) (PTB/LBW) and 278 full-term, normal-weight babies (FTB/NBW). Blood was sampled from the umbilical cord and matched venous blood samples from all neonates between 6-24 hours after delivery. An additional matched venous blood sample was taken from all full-term, normal birth weight newborns between 24-192 hours of life. In both FTB/NBW and PTB/LBW neonates, serum iron decreased 3-fold compared to umbilical blood concentrations within 12h of delivery ( $23.3 \pm 0.35$  vs  $7.5 \pm 0.22$  ng/ml,  $P < 0.001$ ,  $n=425$ ). Hepcidin levels doubled ( $27.0 \pm 0.96$  vs  $52.9 \pm 1.63$  ng/ml,  $P < 0.001$ ,  $n=425$ ). In FTB/NBW neonates, a steady increase in serum iron and TSAT follows (to  $16.5 \pm 3.9 \mu\text{mol/L}$  and  $36.7 \pm 9.2\%$  respectively by 136-192hrs post-delivery), even in the presence of relatively high serum hepcidin levels ( $45.2 \pm 19.1$  ng/ml) suggestive of hepcidin resistance possibly caused by iron saturation of macrophages.

Our findings confirm that a very rapid hypoferraemia occurs in the early hours of post-natal life with evidence that it is mediated by an increase in hepcidin. The strength and consistency of this effect in all neonates indicates that it may have evolved as an innate immune response designed to protect newborns from bacterial septicaemia.

# TABLE OF CONTENTS

STATEMENT OF OWN WORK.....	2
ACKNOWLEDGEMENTS .....	3
ABSTRACT .....	6
TABLE OF CONTENTS .....	7
LIST OF TABLES.....	10
LIST OF FIGURES .....	11
ABBREVIATIONS .....	12
GLOSSARY OF KEY TERMS.....	16
Chapter 1 – Background to Thesis .....	18
1.1 Introduction and Rationale .....	19
1.1.1 Why study neonatal iron metabolism? .....	19
1.1.2 What has previous work by our group provided towards the field?.....	20
1.2 PhD Aims and Objectives .....	22
1.3 PhD Outline .....	23
1.4 Candidate’s Involvement.....	25
1.5 PhD Publications .....	27
1.6 PhD Timeframe.....	30
1.7 Funding.....	30
1.8 Ethics .....	30
1.9 Background .....	31
1.9.1 Ending Preventable Neonatal Mortality .....	31
1.9.2 Neonatal Mortality in Low- and Middle-Income Countries (LMICs).....	32
1.9.3 Neonatal Mortality in The Gambia.....	34
1.9.4 Maternal and Newborn Care at Kanifing General Hospital .....	36
1.9.5 Contribution of Neonatal Infections .....	37
1.9.6 Diagnosis and Treatment of Neonatal Sepsis.....	39
1.9.7 The Threat of Antimicrobial-Resistant Infections .....	41
1.9.8 Interventions to Combat Neonatal Infections .....	42
1.9.9 Transition to Extrauterine Life .....	44
1.9.10 Immune Responses of the Neonate.....	45
1.9.11 Iron in Infection and Immunity .....	47
1.9.12 Dysregulation of Iron Homeostasis .....	50
1.9.13 Pre-Analytical Effects on Hepcidin Measurement.....	51
1.10 References.....	53
Chapter 2 – PhD Study Setting .....	75
2.1 PhD Study Setting.....	76
2.1.1 The Republic of The Gambia .....	76
2.1.2 Kanifing Municipality .....	77
2.1.3 Study Sites: Kanifing General Hospital, The Gambia .....	77
2.1.4 Study Sites: Community Visits (West Coast Region).....	81
2.1.5 Study Sites: Laboratory Work (Kanifing General Hospital and MRCG Keneba Laboratory) .....	82

2.2 References.....	84
Chapter 3 - Hepcidin, serum iron and transferrin saturation in full term and premature infants during the first month of life: A review of existing evidence in humans (Review) .....	85
3.1 ABSTRACT .....	91
3.2 INTRODUCTION .....	92
3.3 METHODS .....	95
3.4 RESULTS .....	97
3.5 DISCUSSION.....	99
3.6 REFERENCES .....	103
3.7 SUPPLEMENTARY MATERIAL .....	141
Chapter 4 - Neonatal iron distribution and infection susceptibility in full term, preterm and low birthweight babies in urban Gambia: study protocol for an observational study (Methodology) .....	142
Chapter 5 – Early postnatal hypoferremia in low birthweight and preterm babies: A prospective cohort study in hospital-delivered Gambian neonates (Main Paper) .....	160
5.1 ABSTRACT .....	167
5.2 INTRODUCTION .....	169
5.3 PARTICIPANTS AND METHODS.....	170
5.4 RESULTS .....	175
5.5 DISCUSSION.....	177
5.6 REFERENCES .....	184
5.7 SUPPLEMENTARY MATERIAL .....	199
Chapter 6 – Iron homeostasis in full term, normal birthweight Gambian neonates over the first week of life (FTB/NBW Paper) .....	215
6.1 ABSTRACT .....	220
6.2 INTRODUCTION .....	221
6.3 SUBJECTS AND METHODS .....	223
6.4 RESULTS .....	227
6.5 DISCUSSION.....	229
6.6 REFERENCES .....	234
Chapter 7 - Discussion .....	257
7.1 A Review of the Neolnnate Study.....	258
7.1.1 Neolnnate Study: Where Did It Begin? (Chapter 1) .....	258
7.1.2 A Review of Iron Homeostasis Over the First Month of Life (Chapter 3) .....	259
7.1.3 Early Postnatal Hypoferremia in Low Birthweight and Preterm Babies (Chapter 5) .....	260
7.1.4 Iron Homeostasis in Full Term, Normal Birthweight Gambian Neonates Over The First Week of Life (Chapter 6) .....	262
7.2 Neolnnate Study: Study Design, Issues Faced and Learning Points for the Future.....	263
7.3 Recommendations For Future Research.....	267
7.3.1 Role of the Placenta in Maternal-Fetal Iron Transfer .....	267

7.3.2 Potential Triggers of Early Postnatal Hypoferremia .....	268
7.3.3 Harnessing Iron to Fight Infections .....	274
7.3.4 Hepcidin Agonists as Hypoferremic Therapies .....	278
7.3.5 Hepcidin-Resistance .....	280
<b>7.4 Conclusions .....</b>	<b>283</b>
<b>7.5 References.....</b>	<b>284</b>
<b>ANNEXES .....</b>	<b>297</b>
<b>ANNEX 1.11 Collaborators and field team details and contributions .....</b>	<b>298</b>
<b>ANNEX 1.12 PhD Timeline.....</b>	<b>301</b>
<b>ANNEX 1.13 MRCG Scientific Coordinating Committee Ethics Letter.....</b>	<b>302</b>
<b>ANNEX 1.14 The Gambia Government/MRC Joint Ethics Committee Letter .....</b>	<b>303</b>
<b>ANNEX 1.15 London School of Hygiene &amp; Tropical Medicine Ethics Letter .....</b>	<b>304</b>
<b>ANNEX 1.16 Oral iron acutely elevates bacterial growth in human serum (FeVir Study).....</b>	<b>305</b>
<b>ANNEX 3.8 New Ballard Score Sheet (Gestational Aging).....</b>	<b>314</b>

## LIST OF TABLES

Table 1.1 Health-related statistics for The Gambia.....	36
Table 1.2 Iron sources of neonatal sepsis causing bacteria.....	48

**Note:** Not including tables in submitted publications.



## LIST OF FIGURES

Figure 1.1 PhD study logo (Neolnnate Study).....	22
Figure 1.2 PhD conceptional framework.....	23
Figure 1.3 Global distribution of newborn death by cause.....	31
Figure 1.4 Distribution of newborn deaths by cause in The Gambia.....	35
Figure 1.5 Overview of innate immune deficiencies of preterm and low birthweight babies..	46
Figure 1.6 The host-pathogen battle of iron.....	48
Figure 1.7 Hepcidin-induced iron homeostasis.....	49
Figure 1.8 The sequestration of iron and its moieties.....	50
Figure 2.1 Map of The Gambia.....	76
Figure 2.2 Map of Kanifing.....	77
Figure 2.3 Kanifing General Hospital Maternity Ward.....	78
Figure 2.4 Community study visits to the homes of recruited newborns.....	81
Figure 2.5 Community study visits involved a complete review of systems.....	82
Figure 2.6 Laboratory bench-top work during PhD study.....	83

**Note:** Not including figures in submitted publications.

## ABBREVIATIONS

AGA	Appropriate for Gestational Age
AGP	Alpha 1-Acid Glycoprotein
ANOVA	Analysis Of Variance
APP	Antimicrobial Protein and Peptides
BCG	Bacillus Calmette–Guérin
BMGF	Bill & Melinda Gates Foundation
BMP6	Bone Morphogenetic Protein 6
CDA	Cord Arterial Blood
CDV	Cord Venous Blood
CI	Confidence Interval
CRE	Carbapenem-Resistant Enterobacteriaceae
CRF	Case Report Form
CRP	C-Reactive Protein
CSF	Cerebrospinal Fluid
CT Scan	Computerized Tomography Scan
CyTOF	Cytometry by Time Of Flight
DC	Dendritic Cells
DFID	Department for International Development
DHS	Demographic and Health Survey
DMT-1	Divalent Metal Transporter 1
ECOWAS	Economic Community of West African States
eCRF	Electronic Case Report Form
EDTA	Ethylenediaminetetraacetic Acid
EFSTH	Edward Francis Small Teaching Hospital
EGF	Epidermal Growth Factor

EIA	Enzyme Immunoassay
ELISA	Enzyme-Linked Immunosorbent Assays
EONS	Early-Onset Neonatal Sepsis
EPO	Erythropoietin
ESBL	Extended Spectrum Beta-Lactamases
FTB	Full-Term Birth
G-CSF	Granulocyte-Colony Stimulating Factor
G6PD	Glucose-6-Phosphate Dehydrogenase
GBS	Group B Streptococci
GCP	Good Clinical Practice
GM-CSF	Granulocyte-Macrophage Colony-Stimulating Factor
HGF	Hepatocyte Growth Factor
HICs	High Income Countries
HIV	Human Immunodeficiency Virus
ID	Identification
IDE	Integrated Development Environment
IGF-1	Insulin-like Growth Factor-1
IgG	Immunoglobulin G
IL-22	Interleukin 22
IL-6	Interleukin 6
IM	Intramuscular
IMCI	Integrated Management of Childhood Illness
IQR	Interquartile Range
IV	Intravenous
KGH	Kanifing General Hospital
KMC	Kangaroo Mother Care
LBW	Low Birth Weight

LMICs	Low- and Middle-Income Countries
LONS	Late-Onset Neonatal Sepsis
LPS	Lipopolysaccharide
LSHTM	London School of Hygiene and Tropical Medicine
MCEE	Maternal Child Epidemiology Estimation
MDA	Mass Drug Administration
MDG	Millennium Development Goal
MHC	Major Histocompatibility Complex
MRC	Medical Research Council
MRCG	Medical Research Council Unit The Gambia at LSHTM
mRNA	Messenger RNA
MRSA	Methicillin-resistant <i>Staphylococcus aureus</i>
MS	Mass Spectrometry
NBS	New Ballard Score
NBW	Normal Birthweight
NDM-1	New Delhi Metallo-Beta-Lactamase 1
NETs	Neutrophil Extracellular Traps
NGAL	Neutrophil Gelatinase-Associated Lipocalin (NGAL)
NGO	Non-Governmental Organizations
NHNES	National Health and Nutrition Examination Survey
NK cells	Natural Killer Cells
OD	Optical Density
PDGF-BB	Platelet-Derived Growth Factor - BB
PhD	Doctor of Philosophy
PI	Principal Investigator
PTB	Preterm Birth
ROS	Reactive Oxygen Species

SBA	Skilled Birth Attendant
SCC	Scientific Coordinating Committee
SD	Standard Deviation
SGA	Small for Gestational Age
siRNA	Small interfering RNA
SOP	Standard Operating Procedure
SSPs	Study Specific Protocols
STAT3	Signal Transducer and Activator of Transcription 3
sTfR	Soluble Transferrin Receptor
TB	Tuberculosis
TFR1	Transferrin Receptor 1
TIBC	Total Iron-Binding Capacity
TLR	Toll-Like Receptor
TSAT	Transferrin Saturation
UIBC	Unbound Iron-Binding Capacity
V1	Venous Bleed 1
V2	Venous Bleed 2
V3	Venous Bleed 3
V4	Venous Bleed 4
VLBW	Very Low Birthweight
VPTB	Very Preterm Birth
WBC	White Blood Cell
WHO	World Health Organisation
YICSS	Young Infants Clinical Signs Studies

## GLOSSARY OF KEY TERMS

<b>Hypoferremia:</b>	A deficiency of iron in the blood.
<b>Nutritional immunity:</b>	To prevent infection from pathogenic organisms by restricting access to essential metals.
<b>Neonatal sepsis:</b>	A type of infection in neonates that specifically refers to the presence of bacteria in the blood stream (due to meningitis, pneumonia, pyelonephritis or gastroenteritis) in the setting of fever. Early-onset (appearing 0-3 days of life) and late-onset (appearing >4 days of life) are further levels of categorisation.
<b>Bacteriostatic:</b>	Substance or process that reversibly inhibits growth or reproduction of bacteria. Different from bactericidal (capable of killing bacteria outright).
<b>Hepcidin:</b>	A 25-amino acid peptide ( <i>HAMP</i> gene) exclusively synthesized by the liver that is the main regulator of systemic iron metabolism. Its primary role is to inhibit iron efflux into the blood by binding to the transmembrane protein, ferroportin.

<b>Transferrin saturation:</b>	The percentage of transferrin molecules that are bound to two iron ions in the ferric form ( $\text{Fe}^{3+}$ ). This is a ratio of serum iron concentration and the total iron-binding capacity (TIBC) expressed as a percentage.
<b>TIBC (total iron-binding capacity):</b>	An indirect measure of the amount of total serum transferrin (apotransferrin, monotransferrin, diferric transferrin) concentration in the circulation.
<b>Post-natal:</b>	The period immediately after the birth of a child.
<b>Preterm birth:</b>	Babies born alive before 37 weeks of pregnancy are completed. Sub-categories included extremely preterm (<28 weeks), very preterm (28-32 weeks) and moderate to late preterm (32-37 weeks).
<b>Low birthweight:</b>	Babies born with a birth weight of <2500g.

# Chapter 1 – Background to Thesis

---

## Summary of Chapter

In this chapter, I briefly introduce the background information and rationale to this thesis. This involves discussing the previous work completed by the HYPO-G Study at MRCG Unit The Gambia by lead investigator Dr Sarah Prentice, which laid the foundations for this further research. I then define the aims, objectives and structure of my research degree. Additionally, I describe the role I played in all aspects of the study, with further information on the research degree timeline. Funding and the subsequent publications produced are also detailed. This chapter then reviews the high-level background literature to this thesis, including research into neonatal mortality linked to infection, the transition to extrauterine life and the role iron plays in infectious disease.



## 1.1 Introduction and Rationale

### 1.1.1 Why study neonatal iron metabolism?

Neonatal sepsis is the third highest cause of death globally, accounting for 225,000 deaths each year.<sup>1</sup> Neonatal infections cause an estimated 23% of all neonatal deaths, with neonatal sepsis alone accounting for 15%. The increasing global threat of antimicrobial resistance will no doubt exacerbate these figures in the future.<sup>2</sup>

Humans undergo the most complex physiological adaption in their life during the transition from a semi-allogeneic, protected foetal setting to a microbe-rich extrauterine environment.<sup>3,4</sup> The initial mass bacterial colonisation of mucosa in the digestive, respiratory, urogenital tracts, as well as the skin<sup>5,6</sup> occurs during the very early neonatal period and can positively affect gut maturation,<sup>7</sup> metabolic homeostasis and immune function<sup>8–10</sup> in early life and beyond. Early-onset neonatal sepsis (EONS) occurs in <72 hours of life, with most causative pathogens being transmitted vertically from mother to infant before or during delivery.<sup>11</sup>

Iron is a cofactor in numerous metabolic pathways that are critical for the human host as well as most pathogens, making it an important mineral in the host-pathogen battle for resources.<sup>12</sup> Therefore, systemic iron distribution is usually strictly regulated.<sup>13</sup> The assimilation of iron from its human host via a plethora of molecular mechanisms (e.g. iron transporters and siderophores<sup>14</sup>), results in increased growth<sup>15</sup> and virulence<sup>16</sup> of many human pathogens. For example, individuals with chronically high iron states (e.g. hemochromatosis), not only have increased free radical redox damage,<sup>17</sup> but an enhanced risk of infection, especially from iron-dependent species of bacteria.<sup>18,19</sup>

The hormone hepcidin is the primary regulator of iron homeostasis.<sup>20</sup> Host inflammatory mediators, IL-6,<sup>21</sup> IL-22<sup>22</sup> and IFN- $\alpha$ ,<sup>23</sup> have been observed to increase the transcription of hepcidin through several Toll-like receptor (TLR) ligands<sup>24</sup> and STAT3 signalling.<sup>25,26</sup> Hepcidin binds to the transmembrane protein ferroportin in macrophages, hepatocytes and enterocytes, resulting in its internalization and degradation.<sup>27–29</sup> Consequently, enteric absorption from dietary iron is reduced, and iron is sequestered in macrophages, which causes a reduction in the extracellular iron concentration. Research carried out in animal models has shown that when the innate immune system is activated, it elicits a hypoferremic response which then limits the risk and severity of bacterial infections.<sup>30–32</sup>

Previous research by our group has suggested that the hepcidin concentration in full-term neonates increases over the first hours of life, which is linked to a significant reduction in serum iron and transferrin saturation over a similar time period.<sup>33</sup> Nevertheless, an accurate and reliable representation of iron homeostasis in all neonates (in particular preterm and/or low birthweight) is still lacking. The process of reviewing this mechanism between multiple studies was previously made harder by the lack of inter-assay standardisation of hepcidin ELISA assays until the recent research conducted by Van der Vorm *et al.*<sup>34</sup>

### **1.1.2 What has previous work by our group provided towards the field?**

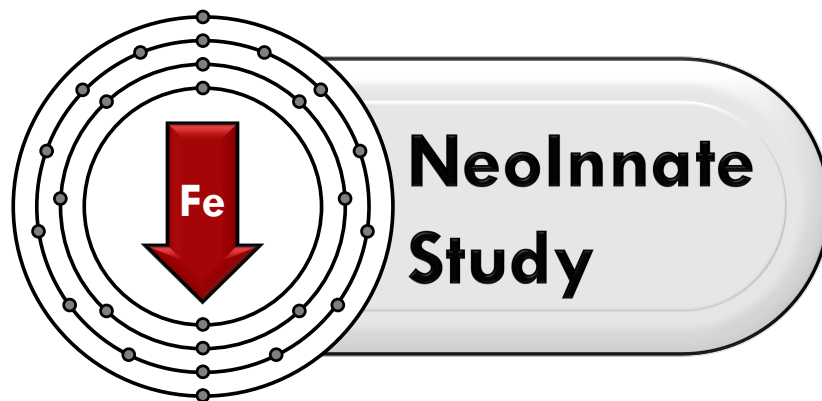
Before the start of this research degree, previous research conducted at MRC Unit The Gambia showed the presence of hepcidin-mediated neonatal hypoferremia in vaginally-delivered healthy full-term neonates in rural Kiang Keneba.<sup>33</sup> This proof-of-concept observational study observed the following key learning points:

- Normal healthy term neonates display a rapid suppression of serum iron and TSAT within the first 6-12 hours post-partum.

- This response is thought to last for 2-3 days, followed by a slow increase up to 92 hours.
- There is a strong negative correlation between hepcidin and serum iron during this period, suggesting that hepcidin regulates this response through the redistribution of iron into circulating macrophages.
- These findings were correlated to levels of IL-6, suggesting that inflammatory stimulation of hepcidin is in part, the driving mechanism of systemic hypoferremia.
- Growth rates of the *ex vivo* micro cultured neonatal sepsis causing bacteria *Escherichia coli*, *Staphylococcus aureus*, *Streptococcus pneumoniae* and *Streptococcus agalactiae* were significantly lower in neonate venous serum (6-24 hr time period) as compared to cord serum.
- Each organism's growth rates was significantly associated with TSAT level in the serum.

To conclude, the evidence to date suggests that neonatal hypoferremia takes place immediately post-partum in healthy term neonates, but there is limited research into the effects of gestational age and birthweight on this method of protection.<sup>33</sup> Our research group pilot data suggests this change in iron distribution could provide a bacteriostatic method of protection to neonates in the first hours of life, when immune defences are immature.<sup>33</sup> The rationale to conduct my research was to confirm this earlier discovery of acute neonatal hypoferremia in full-term neonates in another population, and to test whether preterm (PTB) and/or low birthweight (LBW) infants are equally effective in sequestering iron away from the bloodstream. If this was not the case, we aimed to describe the proportion of such babies that may show ineffective or blunted hypoferremia.

## 1.2 PhD Aims and Objectives



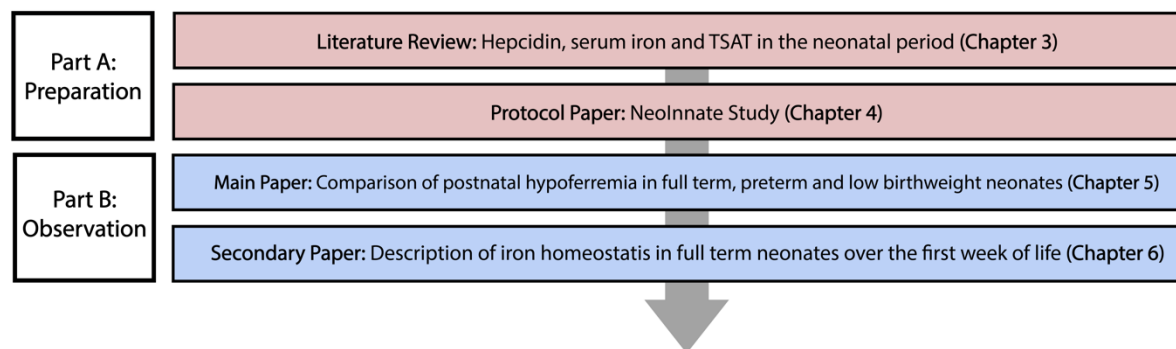
**Figure 1.1** PhD study logo (Neolnnate Study).

**Overall Aim:** To characterise the effects that gestational age and birthweight have on iron distribution immediately after birth and during the first week of life in healthy Gambian neonates.

**Specific Objectives:**

- Conduct a comprehensive review of hepcidin, serum iron and TSAT concentration in neonates in umbilical cord and venous blood (up to 1 month of age).
- Characterise iron metabolism in full-term, preterm and low birthweight neonates at birth and during the first 24 hours of life. Do premature and/or low birthweight neonates have a defect in their ability to sequester iron at 6-24 hours after birth in comparison to full-term newborns with normal birthweight?
- Describe how concentrations of hepcidin, serum iron and TSAT change in full-term neonates (only) at birth and during the first seven days of life.

The PhD conceptual framework is displayed in **Figure 1.2**. This PhD is split into preparation (A) and observation (B) phases. B is itself divided into two sections, with Part B1 focussing on a comparison of iron homeostasis in first hours of life between full-term, normal birthweight (FTB+NBW) neonates and preterm (PTB) and/or low birthweight (LBW) neonates. Part B2 focuses on iron and infection parameters in FTB+NBW neonates after the first week of life only.



**Figure 1.2** PhD conceptual framework.

### 1.3 PhD Outline

This thesis is formatted in a research paper style in accordance with the London School of Hygiene & Tropical Medicine write-up regulations. The manuscripts and published articles are included without any adaption for this thesis. As a result, repetition is common between chapters in regard to materials and methods. This has been reduced with the production of a protocol paper (**Chapter 4**). If supplementary material is part of the manuscripts or publications, it has been added at the end of the chapter. Each chapter begins with a summary page outlining the content of each chapter.

This thesis contains seven chapters, one of which is a published protocol paper (**Chapter 4**), one is a published original research publication (**Chapter 5**) and two submitted manuscripts (**Chapter 3** and **6**). An outline of all the chapters is as follows:

**Chapter 1:** An introduction to the thesis content and structure, with detailed information on the PhD aims, objectives, rationale, candidate involvement, publications and timeframe.

**Chapter 2:** An introduction to the study setting at Kanifing General Hospital and the surrounding communities.

**Chapter 3:** A prepared (submitted) manuscript containing a literature review of the concentrations of hepcidin, serum iron and TSAT in newborn blood in the first weeks of life.

**Chapter 4:** A published research article containing the methodology of all study methods involved in the production of the two original articles (**Chapters 5 & 6**). It should be noted that it was not possible to complete secondary objective (III) in this publication (e.g. *ex vivo* bacterial growth in neonatal serum) due to the lack of adequate blood sample volumes from the initial blood draws.

**Chapter 5:** A published research article comparing the concentrations of hepcidin, serum iron and TSAT in full-term, normal birthweight newborns against preterm and/or low birthweight neonates in the first hours of life.

**Chapter 6:** A prepared (submitted) manuscript describing concentrations of hepcidin, serum iron and TSAT in full term, normal birthweight newborns during the first week of life only.

**Chapter 7:** A final discussion chapter to highlight the overall study findings, limitations of this thesis and reflection on what future research should be conducted in the field of neonatal iron homeostasis and beyond.

## **1.4 Candidate's Involvement**

The original idea of assessing iron and inflammation makers in newborns at birth and shortly afterwards in order to assess if hypoferremia was present came from Dr Sarah Prentice, Dr Carla Cerami (Higher Scientific Officer for MRC Unit The Gambia) and Professor Andrew Prentice (Nutrition Theme Leader for MRC Unit The Gambia). Dr Carla Cerami obtained the funding from the Bill and Melinda Gates Foundation. I assisted in completing the narrative and budget components of the grant (OPP1152353). I worked closely with Dr Carla Cerami (primary supervisor) to develop and implement this observational study and received technical support from Professor Andrew Prentice (secondary supervisor).

As seen in **Chapter 3**, I was responsible for searching the literature, analysing data and formulating the first draft of the narrative review. Dr Carla Cerami and Professor Andrew Prentice (co-authors) supervised me and provided technical support and feedback during the production of the review.

The methodology of the Neolnnate Study (**Chapter 4**) was prepared predominantly from pilot research conducted by Dr Sarah Prentice and Professor Andrew Prentice. Dr Carla Cerami and I adapted previous study procedures and protocols to fit with a change in the study setting. Dr Ousman Jarjou and the nursing team provided clinical insight into the development of study-specific protocols (SPPs). Mr Bakary Sonko oversaw the development of data handling and management, ensuring all aspects followed MRCG data management

policies. Dr Nuradin Ibahim Mohammed provided statistical oversight to ensure the study was adequately powered for our formal analysis.

My role during the Neolnnate Study was a PhD student, along with co-investigator and project manager. I assisted in the writing of the funding application and regular reports to the Bill and Melinda Gates Foundation (BMGF). I assisted in the production of a study budget, monitored spending and procurement/logistics daily. I developed all clinical, data and laboratory study-specific protocols (SSPs). I orchestrated the renovation of a new study-specific laboratory at Kanifing General Hospital (with the help of construction workers, labours and senior hospital staff). I assisted in the production of a successful ethics application to MRCG at LSHTM Scientific Coordinating Committee, MRCG at LSHTM/Gambian Government Joint Ethics Committee and London School of Hygiene and Tropical Medicine Ethics Committee, alongside assisting with the formal registration of the study at ClinicalTrials.gov. I actively supported the communication between Kanifing General Hospital and the Neolnnate study team, holding regular meetings with senior hospital administration. I worked with a team of MRCG at LSHTM staff to arrange a study-specific open day at Kanifing General Hospital, working with hospital administration, local media, international and local non-governmental organisations (NGOs), religious leaders, national and local government and the local population of Kanifing. Furthermore, I aided the assembly of a study team consisting of a research clinician, eight clinical nurses, ten laboratory technicians and two international students by shortlisting candidates, helping conduct interviews and working closely with the Human Resources Department (MRCG at LSHTM). I coordinated the fulfilment of daily project administration and laboratory management (staff appraisal, organising subject community visits, producing staff rotas, and running laboratory controls). I conducted regular shift work at the Neolnnate laboratory, completing primary laboratory analysis and supporting the study nurse team. I developed and verified all electronic data records on the study-specific REDCap (Research Electronic Data Capture) database, executing data curation and management leading to formal



statistical analysis in collaboration with study team members (Dr Carla Cerami and Prof Andrew Prentice) and the study statistician (Dr Nuradin Ibrahim Mohammed). Additionally, I conducted all secondary biochemical processes, sample handling and analysis (ELISA and biochemistry analysis). I provided regular training days to Kanifing General Hospital and Neolnnate Study staff on study procedures, theory and protocols. This work led me to the scientific writing of the manuscripts seen in **Chapters 5 and 6**.

While here, I summarised my contribution to the contents of this thesis, it is essential to note that this study was designed, conducted and analysed in a collaborative approach. I have provided the names and roles of everyone involved in these areas in **Annex 1.11**.

## 1.5 PhD Publications

### Published Papers:

- **Cross JH**, Jarjou O, Mohammed NI, *et al*. Neonatal iron distribution and infection susceptibility in full term, preterm and low birthweight babies in urban Gambia: study protocol for an observational study. *Gates Open Res* 2019;3:1469. (**Chapter 4**)
- **Cross JH**, Jarjou O, Mohammed NI, *et al*. Early postnatal hypoferrremia in low birthweight and preterm babies. *EBioMedicine* 2020;52:102613. (**Chapter 5**)

### Re-Submitted Manuscript:

- **Cross JH**, Prentice AM, Cerami C. Hepcidin, serum iron and transferrin saturation in full term and premature infants during the first month of life: A review of existing evidence in humans. *Current Developments in Nutrition* (re-submitted) (**Chapter 3**)

- **Cross JH**, Jarjou O, Mohammed NI, *et al.* Iron homeostasis in full term, normal birthweight Gambian neonates over the first week of life. *American Journal of Clinical Nutrition* (re-submitted) (**Chapter 6**)

#### **Poster Presentations:**

- Iron and Infection: Neonatal Iron Transition (Neolnnate Study – Prospective). **Cross JH**, Jarjou O, Touray BJB, Prentice AM, Cerami C. Presented at MRC Unit The Gambia 70<sup>th</sup> Anniversary Symposium: 27-29<sup>th</sup> November 2017.
- Iron and Infection: Neonatal Iron Transition (Neolnnate Study – Prospective). **Cross JH**, Jarjou O, Touray BJB, Prentice AM, Cerami C. Presented at MRC Scientific Advisory Board Meeting 2016, MRC Keneba, The Gambia.
- Iron and Infection: Neonatal Iron Transition (Neolnnate Study – Prospective). **Cross JH**, Jarjou O, Touray BJB, Prentice AM, Cerami C. Presented at IUIS-FAIS-ImmunoGambia 2016, MRCG at LSHTM, The Gambia.
- Iron and Infection: Neonatal Iron Transition. **Cross JH**, Prentice AM, Cerami C. Presented at MRC Festival - Show Case, June 2016, MRC Fajara, The Gambia.

#### **Conference Talks:**

- MRC International Nutrition Group. 24<sup>th</sup> April 2019. Neonatal Iron Transition - Neolnnate Study.

- MRC Unit The Gambia – Nutrition Retreat Presentation. 21<sup>st</sup> February 2018. Iron and Infection: Neonatal Iron Transition (Neolnnate Study).
- London School of Hygiene and Tropical Medicine – M. Phil. / Ph.D. Upgrading Seminar. 30<sup>th</sup> September 2017. Iron and Infection: Neonatal Iron Transition (Neolnnate Study).
- MRC Unit The Gambia / MRC International Nutrition Group (Unit Wide). 22<sup>nd</sup> September 2017. Iron and Infection: Neonatal Iron Transition (Neolnnate Study – Prospective).
- MRC Scientific Advisory Board Meeting (10 Minute Elevator Talks). 30<sup>th</sup> March 2016. Iron and Infection: Neonatal Iron Transition (Neolnnate Study – Prospective).
- MRC Unit The Gambia – Nutrition Retreat Presentation. 3<sup>rd</sup> March 2016. Iron and Infection: Neonatal Iron Transition (Neolnnate Study – Prospective).

There was one other paper that I was involved in as part of my laboratory work at MRC Unit The Gambia at LSHTM during my PhD registration. This was in the same field (iron homeostasis) as my thesis, but was unrelated to my PhD research:

- Bah A, Muhammad AK, Wegmüller R, Verhoef H, Goheen MM, Sanyang S, Danso E, Sise EA, Pasricha SRS, Armitage AE, Drakesmith H, **Cross JH**, Moore SE, Cerami C, Prentice AM. Hepcidin-guided screen-and-treat interventions against iron deficiency anaemia in pregnancy: a randomised controlled trial in Gambian women. *Lancet Glob Heal* 2019;**7**:e1564–74

## 1.6 PhD Timeframe

The PhD registration period ran from 28<sup>th</sup> September 2015 until 31<sup>st</sup> December 2019. A Gantt Chart of all PhD and study activities can be observed in **Annex 1.12**.

## 1.7 Funding

The research was undertaken with a research grant provided by the Bill & Melinda Gates Foundation (OPP1152353), under principal investigator Dr Carla Cerami (primary supervisor). The funding agency had no role in the design and conduct of the study and did not have any in the collection, management, analyses or interpretation of the data nor in the preparation, review, or approval of the manuscripts. This research grant was used to pay for all my costs, including my travel and PhD stipend. The Nutrition Theme of the MRC Unit The Gambia at LSHTM are supported by core funding MC-A760-5QX00 to the MRC Unit The Gambia/MRC International Nutrition Group by the UK MRC and the UK Department for the International Development (DFID) under the MRC/DFID Concordat agreement.

## 1.8 Ethics

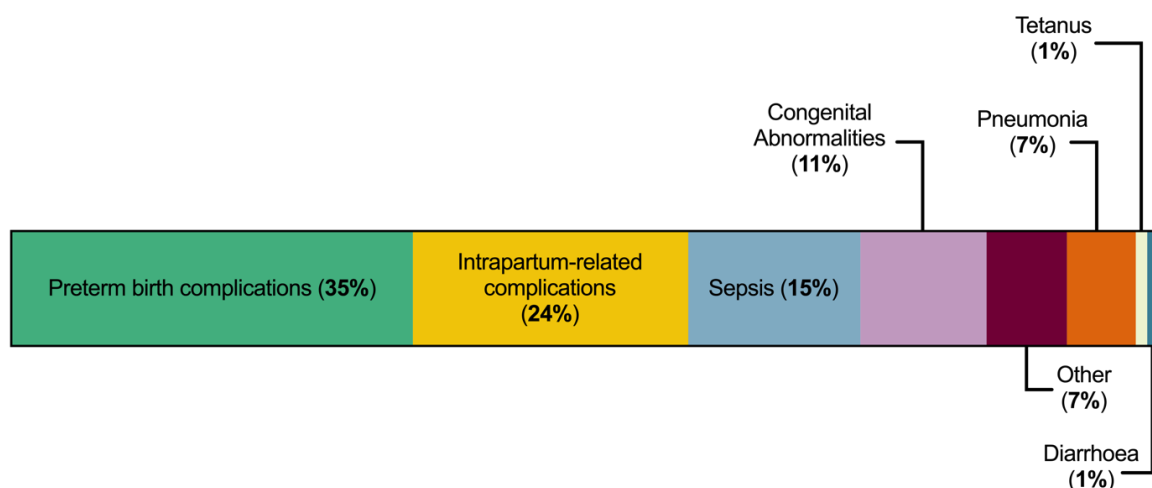
This study has been approved by The Gambia Government/MRC Joint Ethics Committee (no. SCC1525) (**Annex 1.13-1.14**) and Ethics Committee of London School of Hygiene and Tropical Medicine (LSHTM) (ref no. 14316) (**Annex 1.15**). This study was registered with clinicaltrials.gov (NCT03353051) on 27<sup>th</sup> November 2017.

## 1.9 Background

### 1.9.1 Ending Preventable Neonatal Mortality

Since the formation of the World Health Organisation (WHO) Millennium Development Goal 4, significant progress in under-5 child survival has been made over the last two decades.<sup>35</sup> Neonatal deaths have decreased from 5 million (1990) to 2.5 million (2019).<sup>35</sup> Despite this, nearly two million of these newborns die in their first week of life,<sup>36</sup> with 24-45% of all neonatal deaths occurring in the first 24 hours of life.<sup>37</sup> With the substantial reduction in under-5 mortality rates occurring mostly in the older cohort, a more significant proportion of under-5 deaths now occur in the neonatal period. This has been estimated as an increase from 40% in 1990 to 47% globally in 2019.<sup>35</sup>

With the creation of the Sustainable Development Goal 3.2 in 2015, health professionals, academics and policymakers are aiming to end preventable deaths of all children under the age of five by 2030.<sup>38</sup> Part of this is a target for all countries to reduce their neonatal mortality rate to <12 per 1000 live births. Emphasis has also been placed on the ongoing major global health challenge of high rates of stillbirth. It is estimated that a similar number of babies are dying each year before delivery (e.g. 2-3 million).<sup>39</sup>



**Figure 1.3** Global distribution of newborn death by cause. *Source:* Levels & Trends in Child Mortality, WHO and Maternal and Child Epidemiology Estimation Group (MCEE) interim estimates produced in September 2019.<sup>35</sup>

Preterm birth, intrapartum-related complications and infections (including sepsis and pneumonia) are the most common causes of neonatal death.<sup>35</sup> Many of these deaths are due to gaps in the continuum of pre- and post-natal care.<sup>40</sup> This includes pre-pregnancy, antenatal, intrapartum, delivery, postpartum, and postnatal periods for both the mother and her newborn.

### **1.9.2 Neonatal Mortality in Low- and Middle-Income Countries (LMICs)**

Analysis of global neonatal mortality rates has highlighted substantial disparities at a country and regional level. Almost 99% of all neonatal deaths occur in low- and middle-income countries.<sup>37</sup> In particular, the geographical region of Sub-Saharan Africa is struggling to stem the tide, with the highest neonatal mortality rate globally of 28 per 1000 births (2018).<sup>41</sup> This suggests that African newborns are nearly nine times more likely to die in the first month of life than a child born in a high-income country.<sup>41</sup> An extreme example of how much work remains to be done over the next decade is that some regional neonatal death rates in LMICs reach close to 46 deaths per 1000 births.<sup>38</sup>

Simple interventions related to antenatal care, education, nutrition, and maternal health can all help reduce the prevalence of preterm and low birthweight newborns. One of the most effective is the low-cost mass drug distribution of oral iron and folic acid supplementation to pregnant women during pregnancy.<sup>42</sup> This has been shown to reduce the risk of maternal iron deficiency and anaemia at term in the mother, leading to a reduced risk of delivering a preterm and/or low birthweight neonate.

Antenatal care is critical for reducing adverse outcomes in mother and baby. The WHO currently recommend that mothers receive a minimum of eight antenatal care contact visits during their pregnancy.<sup>43</sup> However, only 65% of pregnant mothers in the world achieve four visits before delivery.<sup>44</sup> This is significantly worse in LMICs.<sup>45</sup> Moreover, this analysis does not take into account the skill level of the healthcare provider or the quality of the antenatal care received. The lack of pregnancy-related healthcare education and the high burden of adverse pregnancy and newborn outcomes in LMICs, results in mothers not seeking medical care when necessary. This can apply to both the onset of labour and newborn illness. The lack of regular attendance to antenatal care contact visits can also result in issues relating to micronutrient supplementation, hypertension, immunisations, sexual health screening, and acquiring insecticide-treated mosquito nets. Barriers to these interventions may include physical, financial and cultural aspects.<sup>46</sup>

WHO guidelines support the delivery of a baby by a skilled birth attendant (SBA) at a health facility.<sup>47</sup> Unlike previously, high proportions of deliveries in HICs and LMICs now take place in healthcare facilities.<sup>48,49</sup> The reason for this recommendation was to offer the opportunity to deliver in a safe and clean environment, while providing skilled, good quality essential newborn and maternal care. This policy was designed to allow for the identification of high-risk neonates and promotes their management quickly and effectively. However in LMICs, this has often resulted in more pregnant mothers delivering in unhygienic hospitals with inadequate health infrastructure or technology.<sup>50–52</sup> The upsurge in healthcare facility deliveries has also increased the risk of newborns and mothers acquiring hospital-associated infections, which commonly possess antimicrobial-resistance mechanisms.<sup>53</sup> This is made worse by the fact that many maternity and neonatal wards lack a continuous water supply, basic resuscitation equipment, resources for hand hygiene and waste disposal.<sup>53,54</sup> Similarly, the ratio of skilled healthcare professionals to the number of beds is often far lower than recommended by the WHO.<sup>54</sup> This situation is deteriorating in public hospitals in LMICs faster than anywhere else in the world due to staff attrition to different

countries or to newly established private healthcare facilities.<sup>55</sup> This results in services being overstretched, leading to a reduced quality of care. Mothers as a result may then feel less inclined to remain in the delivery facility for the recommended 24 hours after birth. Ironically, this is the period in which most maternal and neonatal complications present.<sup>56</sup> Studies also suggest that many newborns born in LMICs struggle to receive thermal protection immediately after delivery (e.g. kangaroo mother care (KMC)<sup>57</sup>), hygienic umbilical cord care<sup>58</sup> or early and exclusive breastfeeding.<sup>59</sup> All of these interventions are associated with a reduced risk of infection-related deaths in the newborn.<sup>60–62</sup>

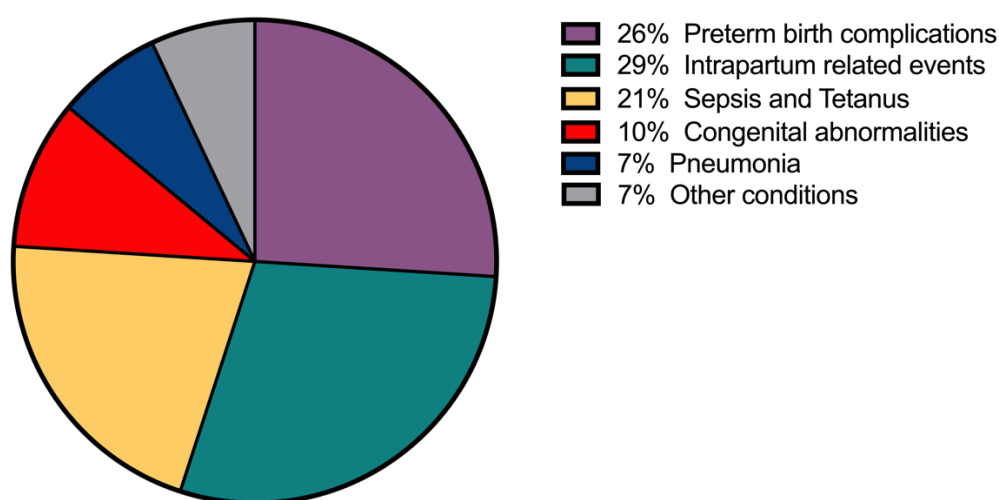
However, high-quality data are required to better understand the fundamental issues affecting newborns and their ability to survive and thrive. Analysis of global mortality rates and their causes has shown that both are influenced by inadequate coverage and poor quality of data.<sup>63</sup> This can result in the misdirection of funds, resulting in inappropriate or weak interventions. Birth registration is a fundamental human right and is vital to economic, social and health planning, yet birth registration is at a prevalence of just 46% in sub-Saharan Africa<sup>64</sup>. The lack of registered births, deaths, and accurate birthweight and delivery records may result in a misunderstanding of the true scale of the issue relating to neonatal mortality. Now there is greater advocacy to collect accurate, timely and disaggregated newborn data from LMICs, which can be used to support evidence-based decision-making, programming and planning.<sup>36,37,65</sup>

### **1.9.3 Neonatal Mortality in The Gambia**

The Gambia has achieved the Millennium Development Goals of improving the under-5 mortality rate, immunisation coverage, the proportion of the population using an improved drinking water source, primary schools enrolment and reduction in malaria disease burden by the 2015 deadline.<sup>66</sup> Much less progress has been made in the reduction of neonatal



mortality rates. National neonatal death levels, as a proportion of under-5 deaths, have increased from 31% in 1990 to 45% in 2017.<sup>67</sup> Like many other LMICs, preterm birth, intrapartum complications and sepsis are responsible for nearly three quarters of all newborn deaths (**Figure 1.4**). On this record, it is unlikely The Gambia can meet its Sustainable Development Goal 3.2 for neonatal mortality by 2030 unless vast and effective interventions are put into practice.



**Figure 1.4** Distribution of newborn deaths by cause in The Gambia. *Source:* Estimates generated by the WHO and Maternal and Child Epidemiology Estimation Group (MCEE) 2018 (<http://data.unicef.org>).<sup>68</sup>

Issues that face neonatal healthcare in The Gambia are similar to those in other LMICs. High levels of poverty, along with a shortage of adequately trained staff and a significant urban bias (66%) in the distribution of the workforce are amongst the issues.<sup>69</sup> In general, Gambian healthcare system suffers from a limited amount of medical supplies and a lack of durable infrastructure and technology.<sup>70</sup> The proportions of birth registration, skilled birth attendants present at delivery, postnatal care for newborns and exclusive breastfeeding proportions show that reasonable improvements can be easily made (**Table 1.1**).

Maternal and Neonatal Health-Related Statistics	Value
Total Births Per Year (000):	81 (2016)
Total Under-5 Population (000):	360 (2016)
Birth Registration (%):	72% (2013)
Total Maternal Deaths Per Year:	590 (2015)
Neonatal Mortality Rate (per 1000 live births)	22 (2013)
Total Stillbirth Rate (per 1000 total births):	24 (2015)
Neonatal Deaths Per Year (as % of all <5 deaths):	45% (2015)
Total Under-5 Deaths Pear Year (000):	5 (2016)
Skilled Birth Attendant (%):	57% (2013)
Postnatal Care for Neonates (%):	6% (2013)
Exclusive Breastfeeding (%):	47% (2013)
Antenatal Care, 4+ Visits (%):	78% (2013)
Low Birthweight Prevalence (%):	10% (2010)
Iron/Folic Acid Supplementation During Pregnancy (%):	45% (2013)
Physician Density (per 1000 population)	0.11 (2015)
Nurse and Midwife Density (per 1000 population)	1.62 (2015)

**Table 1.1** Health-related statistics for The Gambia. *Source:* Countdown to 2030, Maternal, Newborn and Child Survival, Healthy Newborn Network.<sup>67</sup>

#### 1.9.4 Maternal and Newborn Care at Kanifing General Hospital

Annually, Kanifing General Hospital (KGH) provides antenatal care to 500–700 pregnant mothers (unpublished hospital data, 2017), with many presenting very late in their pregnancy. This is thought to be due to pregnant mothers failing to see the advantages gained by early uptake of antenatal care. It is viewed as a curative rather than preventative measure.<sup>71</sup> Local healthcare centres report that few mothers meet the new requirement for eight antenatal visits, with close to half receiving the old requirement of four antenatal visits.<sup>72</sup> This significantly differs from the data produced by a Gambian demographic health survey (DHS).<sup>70</sup>

Antenatal care at Kanifing General Hospital is provided by a group of dedicated, skilled and experienced nurses and midwives. However, as in other hospitals in LMICs, barriers to their care remain. Blood pressure monitoring, urine sampling, blood sampling, tetanus protection, iron supplementation, fetal monitoring and health education are seven routine components of antenatal care that are available at Kanifing General Hospital. Nonetheless, it is rare for all to be available at one antenatal visit. Critical for detecting pre-eclampsia, the lack of back up or charged (i.e. lack of electrical supply) electrical blood pressure monitors results in blood pressure measures being missed. Similarly, due to the lack of full blood count laboratory supplies and broken haemoglobinometers, many weeks' worth of haemoglobin measurements can be lost. This increases the risk of maternal anaemia not being diagnosed, and subsequently could lead to increases rates of stillbirths, low birth weight and preterm babies.<sup>73</sup> During the initiation of this study, it also became apparent that only one Pinard horn for fetal monitoring was available throughout the whole hospital.

The total number of births at Kanifing General Hospital equals close to 3500–4500 per year (based on hospital data from 2014-2015, unpublished). This is 5-6-fold higher than the number receiving antenatal care at the hospital. The percentage of these that are live, low birth weight neonates (<2.5kg) is approximately 10% (based on hospital data from 2014-2015, unpublished). Further details relating to delivery care at Kanifing General Hospital can be seen in **Section 2.1.3**.

### **1.9.5 Contribution of Neonatal Infections**

Neonatal sepsis is the third highest cause of death globally, accounting for 225,000 deaths each year.<sup>1</sup> This results in neonatal infections (e.g. bacterial sepsis, meningitis, pneumonia, and tetanus) causing an estimated 23% of all neonatal deaths. Neonatal sepsis alone accounts for 15% of these deaths,<sup>36</sup> with the majority occurring in Sub-Saharan Africa.<sup>74</sup>

Intrauterine and neonatal infections do not just cause death, but also can result in substantial long-term multisystem morbidity, affecting not just the individual but the local community as well as national productivity.<sup>75,76</sup>

Early-onset neonatal sepsis (EONS) is defined as bacteraemia or bacterial meningitis occurring within <72 hours of life, with most causative pathogens being transmitted vertically from mother to infant before or during delivery.<sup>77</sup> Maternal chorioamnionitis, prematurity, fetal distress (i.e. fetal tachycardia or passage of meconium), prolonged rupture of membranes and maternal colonisation with group B streptococcus (GBS) are all deemed influential risk factors.<sup>77</sup> Quite different to this is late-onset neonatal sepsis, which occurs after this point (>72 hours), with transmission often occurring horizontally from the surrounding delivery environment.<sup>78</sup> As a result, many of the causative microorganisms are nosocomially derived.

In a recent highly informative review conducted by Okomo *et al.*,<sup>79</sup> it is suggested that aetiology data of neonatal infections in sub-Saharan Africa is of poor quality with publications rarely using the STROBE-NI reporting guidelines. Studies that solely observe the epidemiology of neonatal infections in Gambian newborns are lacking. This is despite a high burden of infectious disease in this population and region. In 1999, Mulholland *et al.* suggested that the most important causes of serious, very young infant infections were *Streptococcus pneumoniae*, *Staphylococcus aureus*, and *Salmonella* spp.<sup>80</sup> New data analysis by Okomo *et al.* now suggests that *Staphylococcus aureus*, *Klebsiella* spp. and *Escherichia coli* are the most common causes of bacteraemia and sepsis in African newborns.<sup>79</sup> This change in aetiology over the last twenty years is thought to be due advances in delivery and newborn care along with the introduction of several new vaccines (e.g. Pneumococcal conjugate vaccines and Haemophilus influenza type b).<sup>81</sup> Though not given directly to the newborn, Okomo *et al.* speculates that both vaccines have reduced the transmission of pathogenic isolates in the community. Subsequently, herd immunity could be

affecting the colonisation rates and densities of these and other species of pathogenic and commensal bacteria in the newborn.

Part of the reason why *S. aureus* is found to be a predominant cause of neonatal sepsis in developing countries is believed to be due to its ability to colonise the skin of newborns, mothers and caregivers.<sup>82,83</sup> However, care must be taken with interpreting its apparent high prevalence, as inadequate site sterilisation when blood cultures are taken may also lead to increased rates of false detection.<sup>84</sup> Confirmed *S. aureus* sepsis is strongly associated with higher mortality rates as compared to other bacterial pathogens. This is partly due to metastatic complications in disease progression<sup>85</sup> and its strong association with sepsis in low birthweight newborns.<sup>86</sup>

Enterobacteriaceae such *E. coli* and *Klebsiella* spp. are widespread in the maternal genital tract, and hence neonates acquire these isolates commonly via their gastrointestinal or respiratory tracts during delivery.<sup>87</sup> Okomo *et al.* observed *Klebsiella* spp. to be the second leading cause of neonatal infections in their study.<sup>81</sup> A potential reason for this may be its strong association to nosocomial settings,<sup>88</sup> the capacity to produce biofilms<sup>89</sup> and its ability to display multidrug-resistant phenotypes by producing extended-spectrum  $\beta$ -lactamases (ESBLs) or carbapenemases.<sup>90</sup> With regards to *E. coli*, strains that commonly cause neonatal sepsis often express plasmids, which are essential in processes such as iron acquisition and virulence.<sup>91</sup> Interestingly, *E. coli* and *Klebsiella* strains are associated with risk factors such as prematurity and very low birthweight.<sup>90</sup>

### **1.9.6 Diagnosis and Treatment of Neonatal Sepsis**

Diagnosis of neonatal sepsis remains difficult, partly as there is no accepted definition. Identification with high accuracy and specificity is a challenge not only in developing

countries but even in well-equipped tertiary healthcare facilities in HIC settings.<sup>92</sup> Blood cultures are the gold standard for the diagnosis of neonatal sepsis. Because of economic, logistical and infrastructure-related restrictions, developing countries face multiple challenges when implementing blood culture-based diagnostic testing. Microbiological identification of a pathogen isolated from blood cultures has high specificity, but sensitivity is low. This is supported by the Aetiology of Neonatal Infections in South Asia (ANISA) study,<sup>93</sup> conducted in Bangladesh, India, and Pakistan. This was designed to investigate the incidence of possible severe bacterial infection (pSBI) episodes in the first two months of life and estimate the proportions of bacterial and viral causes. This large study implemented the use of the latest metagenomic approaches, and still only detected 28% of all causative organisms of pSBI. Reasons for low sensitivity of microbiological identification include the collection of small neonatal blood volumes, the presence of low or intermittent isolate concentrations and maternal intrapartum antibiotic exposure.<sup>94</sup> In order to combat this issue, the WHO launched the Integrated Management of Childhood Illness (IMCI) algorithm as part of the WHO Young Infants Clinical Signs Studies (YICSS).<sup>95</sup> This has led to the production of a seven clinical signs diagnosis to use when laboratory services are not available. This clinical algorithm is associated with 85% sensitivity and 75% specificity for severe bacterial infection in the first week of life.<sup>96</sup> Other investigations that are thought to assist in the diagnosis if appropriate include full blood count, chest radiograph, CRP measurement, CSF or skin swabs.<sup>97</sup>

The WHO Integrated Management of Childhood Illness (IMCI) policy suggests that prophylactic intramuscular (IM) or intravenous (IV) ampicillin and gentamicin should be given to neonates with documented risk factors for neonatal sepsis.<sup>98</sup> WHO recommends that this should continue for two days following which the neonate should be reassessed. In LMIC settings, however, the use of parenteral therapies in newborns may be limited by the availability of inpatient neonatal care, transportation to healthcare facilities, economic and

social factors.<sup>99</sup> New interventions to prevent severe neonatal infections are required and are of great interest to the maternal and newborn research community.

### 1.9.7 The Threat of Antimicrobial-Resistant Infections

It has been estimated that the number of people who will die as a result of antibiotic-resistant infections will rise from 700,000 to 10 million each year by 2050.<sup>100</sup> Antibiotic resistance is a major global health threat, with nearly half of the pathogens that cause severe neonatal bacterial infections worldwide reported to be resistant to the first-line (ampicillin or penicillin, and gentamicin) and second-line (third-generation cephalosporins) WHO-recommended treatments.<sup>101</sup> Laxminarayan *et al.* produced estimates suggest 30% of all neonatal deaths can be attributable to multidrug-resistant pathogens.<sup>102</sup> A disproportionate number of these deaths occur in developing countries, and the rates are increasing. In particular, a small group of studies conducted in Sub-Saharan Africa show a high prevalence of resistance to recommended empirical therapies to neonatal sepsis.<sup>103,104</sup>

In developing countries, gram-negative bacteria have frequently been reported as the cause of neonatal sepsis.<sup>79</sup> This is a worrying trend, with many gram-negative bacteria intrinsically resistant to many antibiotics, along with the ability to transfer genetic material of new resistance mechanisms between species.<sup>105</sup> *Klebsiella* spp. and *E. coli* can both carry extended-spectrum beta-lactamase (ESBL) producing plasmids, enabling them to become resistant to a wide variety of penicillin and cephalosporin antibiotics.<sup>106</sup> In this scenario, carbapenems are the last remaining treatment option,<sup>106</sup> however, the availability of second-line antibiotics in developing countries is known to be poor.<sup>107</sup> *K. pneumoniae* is also the leading cause of infections caused by carbapenem-resistant bacteria worldwide.<sup>108</sup> The acquisition of an enzyme called New Delhi metallo-beta-lactamase (NDM-1) is a significant reason for this.<sup>109</sup> Bacteria with this genetic element can cause disease with increased

morbidity and mortality,<sup>110</sup> especially in preterm newborns.<sup>111</sup> Previous authors have commented that strains harbouring these antimicrobial-resistant components are a massive threat to neonatal units, particularly in LMIC hospitals as neonates are subjected to poor infection control and high antimicrobial exposure.<sup>112</sup>

With regards to *S. aureus* antimicrobial resistance, methicillin-resistant *Staphylococcus aureus* (MRSA) infections have now spread worldwide,<sup>113</sup> with some of the fastest rates of the increase occurring in community settings.<sup>114</sup> The WHO now reports that in some locations of Africa, 80% of *S. aureus* infection are caused by MRSA.<sup>115</sup> Nevertheless, optimism remains, with MRSA still remaining sensitive to glycopeptides such as vancomycin and teicoplanin, and other antibiotics like linezolid, tigecycline, and daptomycin.<sup>114</sup> However, many of these drug options are expensive, regularly unavailable in LMICs and have potent side-effects on newborns.<sup>98</sup>

### **1.9.8 Interventions to Combat Neonatal Infections**

*Staphylococcus aureus*, *Klebsiella* spp., and *Escherichia coli* are all pathogens associated with poor hygiene in hospital environments, suggesting that new interventions are required in sub-Saharan hospitals to increase infection control from surfaces, hands and water. One such intervention is the use of a bundle of hygiene-based interventions for a hospital maternity ward. A study that implemented such a bundle provided enhanced infection prevention and control training, clean delivery kits (e.g. a blade, cord thread, antiseptics), text message reminders, alcohol hand rub, timed environmental cleaning and the introduction of bathing babies >1.5 kg with 2% chlorhexidine gluconate.<sup>116</sup> This study subsequently observed reductions in sepsis and death rates in hospitalised neonates in a high-risk setting in a developing country. These kinds of interventions offer simplicity and



reasonable levels of sustainability, with the production of standardised protocols helping to improve aseptic practices in the delivery rooms and neonatal units.

A further intervention toward the prevention of neonatal sepsis in LMICs is the use of oral azithromycin given to the mother during labour.<sup>117</sup> Azithromycin is a macrolide antibiotic that has a broad antimicrobial activity. Azithromycin has also been found to reduce the proportion of low birthweight births by a quarter, when given intrapartum to the mother for the last few months of pregnancy.<sup>118</sup> Equally, it has been able to reduce overall under-5 mortality when given in the form of a mass drug administration trial.<sup>119</sup> More recent studies have shown that azithromycin during labour can significantly reduce bacterial carriage (e.g. mainly *S. aureus*, *GBS* and *S. pneumoniae*) both in the mother and her newborn.<sup>117</sup> This suggests that it halts the vertical transmission of neonatal sepsis-causing organisms from mother to child and hence may lower their risk of neonatal sepsis, pneumonia and meningitis. It is now believed that this simple one dose intervention could dramatically improve neonatal mortality in LMICs. However, questions remain about the effect MDA of antibiotics will have on the levels of antimicrobial resistance in the mother, newborn and the broader community.<sup>120</sup>

Symbiotics are a combination of probiotics and prebiotics; the latter is added to promote growth and sustain the colonisation of the probiotic bacterial strain. Studies looking at the effects of symbiotics, prebiotics and probiotics have grown more popular over recent years. This is primarily due to the naive immune system of the neonate and the undeveloped gastrointestinal microbiome making it easier to establish probiotic strains.<sup>121</sup> One of the main successes of this area of study is the use of probiotics to reduce the incidence and severity of necrotising enterocolitis in premature infants.<sup>122,123</sup> Now, research suggests that symbiotics can be used to reduce neonatal sepsis mortality rates in LMICs.<sup>124,125</sup> One of the most prominent studies using symbiotics, is the one conducted by Panigrahi *et al.*<sup>126</sup> This was a randomised, double-blind, placebo-controlled oral symbiotic trial of *Lactobacillus plantarum* plus fructo-oligosaccharide in India. This study identified that a single symbiotic preparation was protective against neonatal sepsis and death in the first week of life. This is

an exciting finding, that may relate to the findings of our study, as *Lactobacillus plantarum* is an organism that does not require iron for growth, and instead uses manganese in many of its metalloenzymes.<sup>127</sup> The establishment of *Lactobacillus plantarum* colonisation at birth could be out competing pathogenic species (i.e. that require iron) for space and niches, due to the hypoferremic conditions. Panigrahi *et al.* believe that these findings suggest that this cost-effective method could be used in developing countries to reduce the rates of neonatal sepsis.<sup>126</sup>

### 1.9.9 Transition to Extrauterine Life

The higher rates of mortality relating to severe bacterial infections in newborns are suggested to be partially due to the immense physiological and immunological strain on the neonate in the first few days of life. The transition from fetal to extrauterine life leads to all organ systems being involved in major instantaneous adaptations to respiratory exchange, cardiovascular flow, endocrine function, haematological maturation, substrate metabolism and thermogenesis.<sup>128</sup>

Though the sterile womb paradigm versus *in utero* colonisation hypothesis continues to divide,<sup>129,130</sup> childbirth is still characterised by a neonate descending from a semi-allogeneic, protected environment to one that is far more abundant in a diverse array of microbes. This is the initial focal point of mass bacterial colonisation of the skin and gastrointestinal tract of the neonate, resulting in changes to nutritional and immunological functions in early life and beyond.<sup>5,6</sup>

Regarding iron homeostasis, as seen in our study, increases in newborn haematocrit and haemoglobin occur within the first hours post-delivery. This is likely due to post-delivery dehydration, as well as vasomotor instability and venous pooling.<sup>131</sup> Similarly, serum ferritin

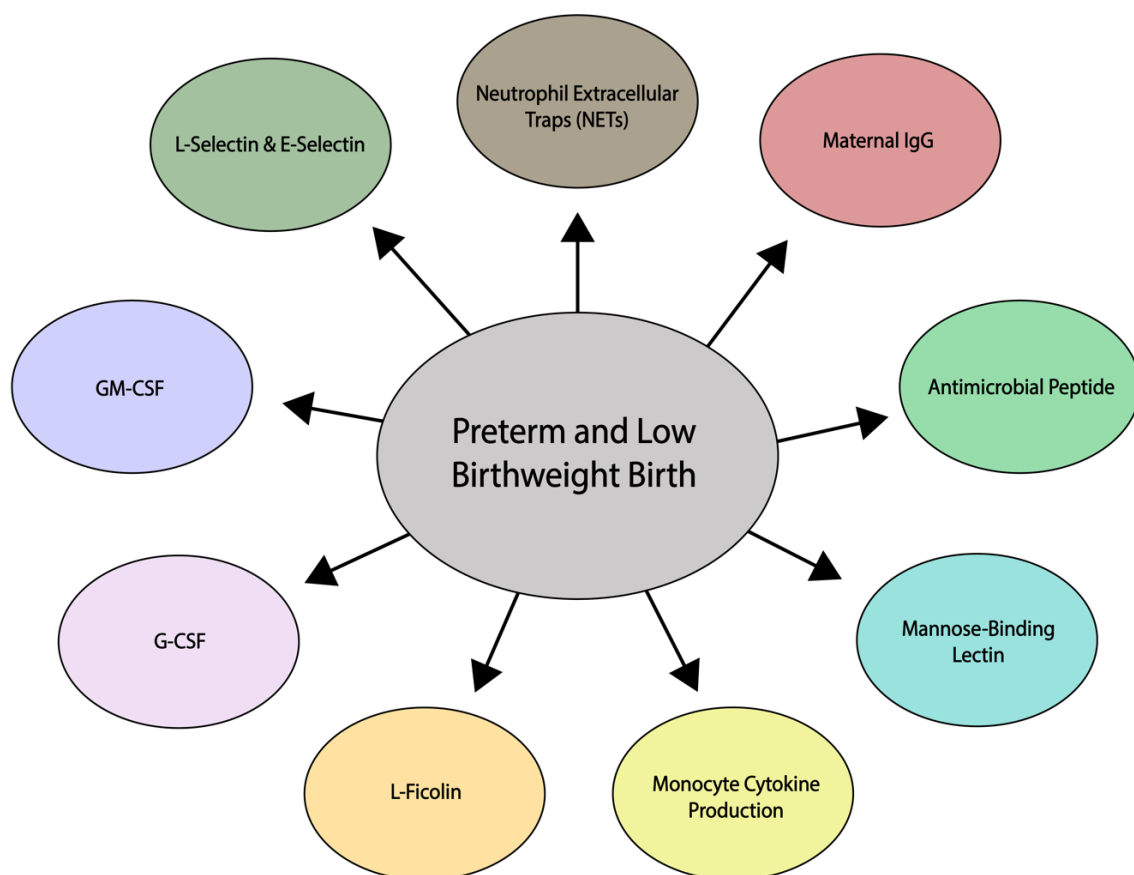
levels increase due to the physiological haemolysis of fetal red blood cells, which contain ferritin in high concentrations.<sup>132</sup> This process leads to fetal haemoglobin recycling into the adult form to aid respiratory exchange in the extrauterine environment.<sup>133</sup> A frequent side-effect of this red cell turnover is hyperbilirubinemia (i.e. physiological or idiopathic jaundice), which is a common clinical issue in most (60-80%) neonates in the first weeks of life.<sup>134</sup>

### 1.9.10 Immune Responses of the Neonate

Neonates are also partly hindered by possessing a functionally immature and developing immune system. The broad-range killing capacity of the innate immune system consists of granulocytes, antigen-presenting cells, NK cells and  $\gamma\delta$ -T cells.<sup>135</sup> Neutrophils are the predominant cell type of the innate immune system, with their role being to engulf and degrade pathogens during infection. Nevertheless, neonatal neutrophils at birth can be measured as low as  $1.5 \times 10^9$  cells/L blood, compared to  $4.4 \times 10^9$  cells/L in adults.<sup>136</sup> Neonatal neutrophils also possess qualitative deficiencies in respect to low levels of expression of TLR4 (critical for IL-6 expression), cell surface L-selectin, Mac-1 (critical for transmigration) and neutrophil extracellular traps (NETs).<sup>137,138</sup> These deficiencies are even more pronounced in preterm neonates. Similarly, DC cells,  $\gamma\delta$ -T cells and monocytes are also all in low concentrations in the neonate and exhibit low expression levels of MHC-II, CD80, CD86, CD40 and ICAM-115. This is thought to inhibit their ability to activate antigen-specific B and T cell populations.<sup>136</sup>

The immune system of premature neonates is further underdeveloped when compared to full-term newborns (**Figure 1.5**). Antimicrobial proteins and peptides (APPs) are released to destroy pathogens by iron-binding, enzymatic destruction, zinc deprivation and membrane pore formation. Such proteins and peptides are secreted into the vernix caseosa found coating the skin, respiratory and gastrointestinal tracts of newborn babies.<sup>139</sup> However, the

production of APPs and vernix caseosa is positively correlated with gestational age,<sup>140</sup> potentially leading to early preterms being at higher risk of infection. Preterms also start life with a lower endowment of antigen-specific IgG than full-term neonates, resulting in reduced opsonisation and phagocytosis. This is due to IgG being transferred across the placenta from the maternal circulation in large amounts after 32 weeks of gestation.<sup>140</sup> In full-term neonates, the complement pathways (classical, alternative and lectin) are generally similar to adult levels, this aids the destruction of bacterial cells. However, these pathways are also all immature in preterm neonates.<sup>141</sup>



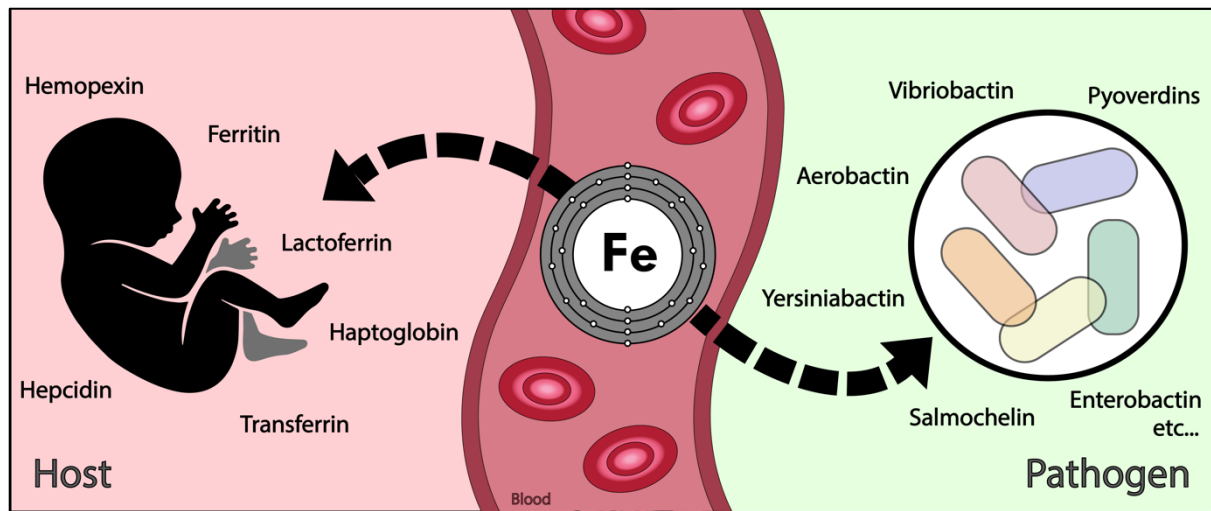
**Figure 1.5** An overview of innate immune deficiencies of preterm and low birthweight babies. Adapted from Melville *et al.*<sup>140</sup>

Immunological deficiencies in neonates (exaggerated in low birthweight and preterm babies) no doubt reflect in the typical clinical presentation of neonatal sepsis. This presentation is

often rapid in progression, with many nonspecific clinical signs.<sup>97</sup> This is further confirmed by unusual haematological laboratory results coming from newborn blood samples.<sup>142</sup> Our study experienced such results, with even healthy newborns found to have increased white blood cell counts from cord and venous samples; though we suspect that this might be an artefact of nucleated red blood cells being identified as granulocytes.

### **1.9.11 Iron in Infection and Immunity**

Iron is a cofactor in numerous metabolic pathways that are critical for the human host. Transition metals (e.g. Fe, Cu and Mg) are commonly built-in to metalloenzymes, storage proteins and transcription factors.<sup>143</sup> This results in children and pregnant woman requiring adequate iron stores for growth and development. The lack of such stores leads to iron deficiency and anaemia, which subsequently negatively affects the immune response to infection.<sup>144</sup> However, the catalytic action of these metals also potentiates their toxicity when in high concentrations (i.e. equalling tissue damage and inflammation), so the levels of these metals must be well regulated.<sup>145</sup> The mechanisms that regulate the toxicity of free transition metals in the body, additionally function as a countermeasure against local pathogens (**Figure 1.6**).<sup>143</sup> This is seen clinically with individuals suffering from high iron states (e.g. haemochromatosis), not only resulting in free radical redox damage but also an enhanced risk of infection from iron-dependent species of bacteria.<sup>18,19</sup> This host-pathogen battle for resources is due to iron also being a vital commodity for growth, replication and virulence of most pathogenic microorganisms. Subsequently, pathogens have evolved counteractions to the host's iron-restriction mechanisms and assimilate iron from their human host.<sup>146</sup>



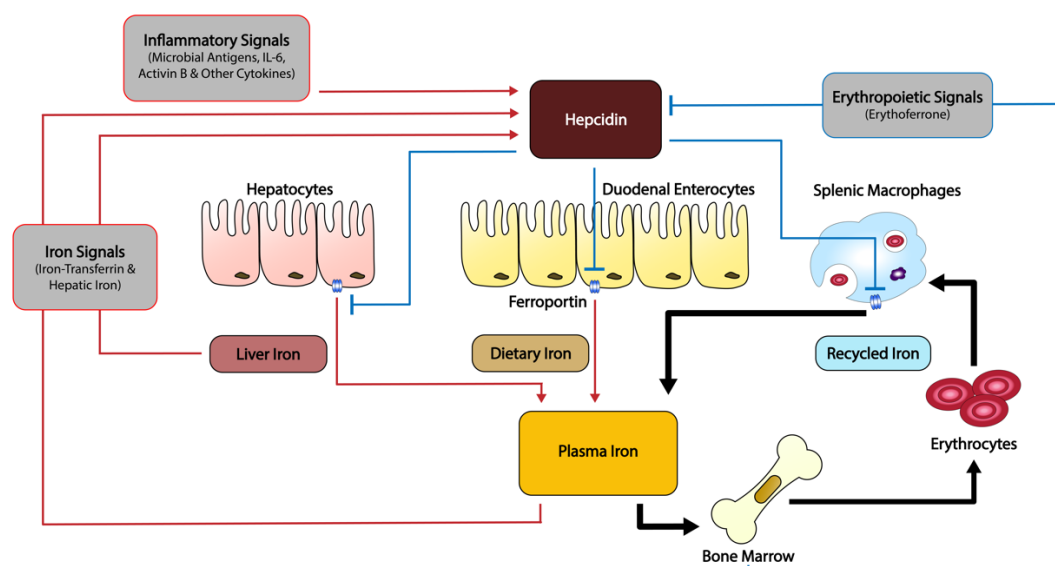
**Figure 1.6** The host-pathogen battle of iron.

Neonatal sepsis-causing bacterial pathogens can produce a plethora of molecular mechanisms such as siderophores and iron-specific channels (**Table 1.2**). These iron acquisition genes are frequently concentrated on high pathogenicity islands.<sup>147</sup>

Substrate	<i>Staphylococcus aureus</i>	<i>Escherichia coli</i>	<i>Streptococcus pneumoniae</i>	<i>Klebsiella pneumoniae</i>	<i>Streptococcus agalactiae</i>
Haem	YES 148,149,150	YES 151,152,153	YES 154,155,156	?	YES 157
Haemoglobin	YES 158,159,160	YES 151,152,161	YES 154,155,156	YES 147	?
Haem-Haemopexin	NO 159	?	?	?	?
Haemoglobin-Haptoglobin	YES 162,163	?	?	YES 147	?
Transferrin	YES 165,166	YES 146,167,168,169	NO 154,155,170	YES 171,172,173,174	?
Direct Transferrin	YES 176,177	? <sup>178</sup>	?	?	?
Lactoferrin	? 179,180	? 178	NO 154,155,170	YES 147	?
Ferric Iron	YES 181,146	YES 182	YES 183,184	YES 171,172,173	?
Ferrous Iron	? 185	YES 186,187,188	YES 183	YES 189,190	YES 191
Ferritin	?	YES 192	?	?	?
NTBI	YES 193	YES 194,195	?	?	?

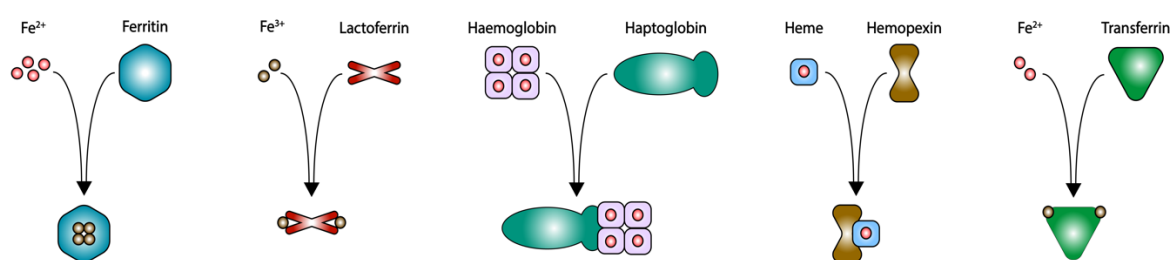
**Table 1.2** Iron sources of neonatal sepsis causing bacteria. *Source:* Adapted from Andrews *et al.*<sup>196</sup>

In 2000-2001, three independent research groups discovered the hormone hepcidin.<sup>197–199</sup> Hepcidin is now understood to be the master regulator of human iron homeostasis, and consequently iron-restriction.<sup>200</sup> Host inflammatory mediators, IL-6<sup>21</sup>, IL-22<sup>22</sup> and Type 1 interferon<sup>23</sup> increase transcription of hepcidin in the liver through several toll-like receptor (TLR) ligands<sup>24</sup> and STAT3 signalling.<sup>24,25</sup> This results in the internalization of the transmembrane protein ferroportin in macrophages and enterocytes. This leads to the breakdown of ferroportin in the lysosome. Consequently, enteric absorption of dietary iron is reduced, and sequestration of iron in macrophages causes a decrease in systemic iron concentration in the circulation.<sup>201</sup> In neonates, due to the relatively small amounts of iron absorbed enterically, iron restriction mainly takes place in circulating macrophages. Hepcidin synthesis is also known to be controlled by high iron stores, hypoxia, and erythropoiesis (Figure 1.7).<sup>200</sup>



**Figure 1.7** Hepcidin-induced iron homeostasis. *Source:* Adapted from Ganz *et al.*<sup>200</sup>

The human host also produces many host chaperone molecules such as transferrin, lactoferrin, haptoglobin, hemopexin (**Figure 1.8**) to limit the acquisition of iron and its moieties (haem and haemoglobin), thus making iron harder to sequester from the human host.<sup>202</sup>



**Figure 1.8** The sequestration of iron and its moieties. Source: Adapted from Parrow *et al.*<sup>202</sup>

### 1.9.12 Dysregulation of Iron Homeostasis

For the reasons stated above, iron homeostasis needs to be maintained at all times. Iron supplementation is widely regarded as one of the most critical public health interventions in low and middle income countries, reducing the morbidity and mortality linked to iron deficiency and anaemia.<sup>42</sup> However, reports have suggested that iron supplementation in areas with high burdens of infectious disease can increase the risk of severe bacterial morbidity and mortality.<sup>203</sup> We suggest that a study conducted by our team (and led by the candidate) partly explains the mechanism. In the previous FeVir study,<sup>204</sup> we collected the blood of 48 adult male Gambian subjects immediately before and four hours after taking 400 mg ferrous sulphate orally. We then performed full blood counts and analysis of serum biochemical parameters, including serum iron, transferrin, ferritin, and transferrin saturation (TSAT). Additionally, we inoculated these paired serum samples with different sepsis causing organisms such as *Staphylococcus aureus*, *Staphylococcus epidermidis*, *Salmonella enterica* serovar Typhimurium and *Escherichia coli* in an *ex vivo* bacterial growth assay. This study concluded that *ex vivo* bacterial growth over twelve hours (36h for *Y.*



*enterocolitica*) was significantly increased by supplemental iron and strongly correlated with TSAT and serum iron concentration (**Annex 1.16**).

A clinical study by Barry *et al.* showed similar results in a different context.<sup>205</sup> Barry *et al.* discovered that Polynesian neonates were twenty-fold more likely to have neonatal sepsis compared to European neonates. This research uncovered that this was due to intramuscular iron dextran injections given to the neonate if they were thought to be anaemic. Barry *et al.* also described that there was not only a difference in the incidence of sepsis and mortality between those given iron dextran injections but additionally, a difference in the type of causative organism. Further data showed that iron dextran had impaired the nutritional immunity of the newborns and changed the causative organism from *S. aureus* to *E. coli*. This finding associates closely with the differences in growth seen between *S. aureus* and *E. coli* isolates in response to TSAT level seen in our previous FeVir study.<sup>204</sup>

Both the *ex vivo* and *in vivo* studies suggest that lower iron concentrations in the blood are vital to limit the growth of bacteria and severity of sepsis caused by common neonatal sepsis-causing pathogens.

### **1.9.13 Pre-Analytical Effects on Hepcidin Measurement**

As hepcidin is the master regulator of iron homeostasis, its measurement can be crucial to the diagnosis of several iron-related disorders. However, its accurate measurement is complicated. Hepcidin is at the centre of the iron, infection and inflammation axis.<sup>206</sup> Significant increases in hepcidin concentration occur in response to iron administration and/or inflammatory stimuli. Subsequently, this can cause several clinical conditions and treatments to confound systemic hepcidin measurements. This is observed for chronic

kidney disease (increase),<sup>207</sup> erythrocyte transfusions (increase),<sup>208</sup> alcohol abuse (decrease)<sup>209</sup> and chronic hepatitis C viral infection (decrease)<sup>210</sup> all having an effect. These observations result in the need for a detailed medical history to be collected when researching the molecular mechanisms related to iron homeostasis in the human host.

Serum hepcidin concentrations also vary significantly between healthy subjects, as evidenced by the published reference ranges.<sup>211,212</sup> Itkonen *et al.* noted that reference values for adult females (18–50 yrs) ranged between 0.4–9.2 nmol/L, 0.7–16.8 nmol/L for females >50 yrs and for adult males (≥ 18 yrs) between 1.1–15.6 nmol/L.<sup>212</sup> These results suggest lower hepcidin levels are observed in pre-menopausal women, as well as a significant difference between sexes (i.e. higher concentrations in men). In-subject variation is also detected, with fasting and diurnal rhythm both found to play a role.<sup>213,214</sup> This is seen with serum hepcidin concentrations being found to be lowest in the early morning, increase over the day, followed by a return to more moderate levels by evening.

Further complications of measuring hepcidin concentration in human urine, serum and plasma come from laboratory-based issues. Of the two main methods of measuring hepcidin, mass spectrometry-based methods (MS) are expensive but are accurate, more reproducible and measure only the native complete form of hepcidin, hepcidin-25.<sup>215</sup> Enzyme-linked immunosorbent assay-based methods (ELISA) are more cost-effective; however, these methods additionally measure the other isoforms of hepcidin, hepcidin-20 and hepcidin-22.<sup>215</sup> The effect of these different test attributes on the clinical diagnosis of iron-related disorders is still unclear. The measurement of hepcidin by ELISA-based methods is further hindered by hepcidin being a relatively small size (i.e. 25 amino acids) with few antigenic epitopes.<sup>216</sup> Antibody generation required for the production of the ELISA-based method is hampered by the high degree of genetic conservation between animal species, making it a challenge to produce viable antibodies from exposure to the protein in

the host animal.<sup>217</sup> Production of synthetic alternatives is also challenging due to hepcidin's complex hairpin-like conformation with four disulphide bonds.<sup>218</sup>

An essential aspect of the laboratory handling process of samples requiring serum hepcidin measurement is the monitoring of freeze-thaw cycles. Serum hepcidin is reported to only remain stable for two to four freeze-thaw cycles, dependent on the analysis method.<sup>219,220</sup> Additionally, Itkonen *et al.* has noted that hepcidin concentration starts to decrease after 24-48 hours storage at room temperature, six days at 4°C, 42 days at -20°C, and over two years at -80°C.<sup>212</sup> This is not bettered by hepcidin's ability to aggregate and adhere to plastic laboratory test tubes and pipette tips.<sup>221</sup>

Previously, the analysis of hepcidin concentration has been hindered by the lack of a commutable calibrator, a reference material, and a reference method.<sup>221</sup> Subsequently, analysis of serum hepcidin levels from various publications share correlation; however, absolute values regularly vary considerably between the ELISA test kits used (**Chapter 3**) This is observed with low values being associated with the use of DRG ELISA kits, and high values linked with the use of Intrinsic Lifesciences and Bachem ELISA kits. To combat this Van der Vorm *et al.* produced a commutable candidate reference material that was a native, lyophilized plasma with cryoprotectant.<sup>34</sup> This material was refined and is now freely available for purchase.<sup>222</sup> However, this product was not available at the time of laboratory analysis during this study.

## 1.10 References

- 1 GBD 2016 Causes of Death Collaborators M, Abajobir AA, Abbafati C, *et al.* Global, regional, and national age-sex specific mortality for 264 causes of death, 1980-2016: a systematic analysis for the Global Burden of Disease Study 2016. *Lancet* 2017;

- 390:** 1151–210.
- 2 Laxminarayan R, Duse A, Wattal C, *et al.* Antibiotic resistance—the need for global solutions. *Lancet Infect Dis* 2013; **13**: 1057–98.
  - 3 de Goffau MC, Lager S, Sovio U, *et al.* Human placenta has no microbiome but can contain potential pathogens. *Nature* 2019; **572**: 329–34.
  - 4 Rutayisire E, Huang K, Liu Y, Tao F. The mode of delivery affects the diversity and colonization pattern of the gut microbiota during the first year of infants' life: a systematic review. *BMC Gastroenterol* 2016; **16**: 86.
  - 5 Dominguez-Bello MG, Costello EK, Contreras M, *et al.* Delivery mode shapes the acquisition and structure of the initial microbiota across multiple body habitats in newborns. *Proc Natl Acad Sci* 2010; **107**: 11971–5.
  - 6 Houghteling PD, Walker WA. Why Is Initial Bacterial Colonization of the Intestine Important to Infants' and Children's Health? *J Pediatr Gastroenterol Nutr* 2015; **60**: 294–307.
  - 7 Tourneur E, Chassin C. Neonatal Immune Adaptation of the Gut and Its Role during Infections. *J Immunol Res* 2013; **2013**: e270301.
  - 8 West CE, Jenmalm MC, Prescott SL. The gut microbiota and its role in the development of allergic disease: a wider perspective. *Clin Exp Allergy* 2015; **45**: 43–53.
  - 9 Noverr MC, Falkowski NR, McDonald RA, McKenzie AN, Huffnagle GB. Development of allergic airway disease in mice following antibiotic therapy and fungal microbiota increase: role of host genetics, antigen, and interleukin-13. *Infect Immun* 2005; **73**: 30–8.
  - 10 Wopereis H, Oozeer R, Knipping K, Belzer C, Knol J. The first thousand days - intestinal microbiology of early life: establishing a symbiosis. *Pediatr Allergy Immunol* 2014; **25**: 428–38.
  - 11 Chan GJ, Lee AC, Baqui AH, Tan J, Black RE. Prevalence of early-onset neonatal infection among newborns of mothers with bacterial infection or colonization: a

- systematic review and meta-analysis. *BMC Infect Dis* 2015; **15**: 118.
- 12 Schaible UE, Kaufmann SHE. Iron and microbial infection. *Nat Rev Microbiol* 2004; **2**: 946–53.
  - 13 Pigeon C, Ilyin G, Courselaud B, *et al.* A New Mouse Liver-specific Gene, Encoding a Protein Homologous to Human Antimicrobial Peptide Hepcidin, Is Overexpressed during Iron Overload. *J Biol Chem* 2001; **276**: 7811–9.
  - 14 Caza M, Kronstad JW. Shared and distinct mechanisms of iron acquisition by bacterial and fungal pathogens of humans. *Front Cell Infect Microbiol* 2013; **3**:00080.
  - 15 Cross JH, Bradbury RS, Fulford AJ, *et al.* Oral iron acutely elevates bacterial growth in human serum. *Sci Rep* 2015; **5**: e16670.
  - 16 Litwin CM, Calderwood SB. Role of iron in refutation of virulence genes. *Clin Microbiol Rev* 1993; **6**: 137–49.
  - 17 Pietrangelo A. Hereditary hemochromatosis. *Biochim Biophys Acta - Mol Cell Res* 2006; **1763**: 700–10.
  - 18 Frank KM, Schneewind O, Shieh W-J. Investigation of a Researcher's Death Due to Septicemic Plague. *N Engl J Med* 2011; **364**: 2563–4.
  - 19 Khan FA, Fisher MA, Khakoo RA. Association of hemochromatosis with infectious diseases: expanding spectrum. *Int J Infect Dis* 2007; **11**: 482–7.
  - 20 Collins JF, Wessling-Resnick M, Knutson MD. Hepcidin regulation of iron transport. *J Nutr* 2008; **138**: 2284–8.
  - 21 Nemeth E, Rivera S, Gabayan V, *et al.* IL-6 mediates hypoferremia of inflammation by inducing the synthesis of the iron regulatory hormone hepcidin. *J Clin Invest* 2004; **113**: 1271–6.
  - 22 Armitage AE, Eddowes LA, Gileadi U, *et al.* Hepcidin regulation by innate immune and infectious stimuli. *Blood* 2011; **118**: 4129–39.
  - 23 Ryan JD, Altamura S, Devitt E, *et al.* Pegylated interferon- $\alpha$  induced hypoferremia is associated with the immediate response to treatment in hepatitis C. *Hepatology* 2012; **56**: 492–500.

- 24 Peyssonnaud C, Zinkernagel AS, Datta V, Lauth X, Johnson RS, Nizet V. TLR4-dependent hepcidin expression by myeloid cells in response to bacterial pathogens. *Blood* 2006; **107**: 3727–32.
- 25 Verga Falzacappa M V., Vujic Spasic M, Kessler R, Stolte J, Hentze MW, Muckenthaler MU. STAT3 mediates hepatic hepcidin expression and its inflammatory stimulation. *Blood* 2007; **109**: 353–8.
- 26 Wrighting DM, Andrews NC. Interleukin-6 induces hepcidin expression through STAT3. *Blood* 2006; **108**: 3204–9.
- 27 Delaby C, Rondeau C, Pouzet CC, *et al.* Subcellular Localization of Iron and Heme Metabolism Related Proteins at Early Stages of Erythrophagocytosis. *PLoS One* 2012; **7**: e42199.
- 28 Chaston T, Chung B, Mascarenhas M, *et al.* Evidence for differential effects of hepcidin in macrophages and intestinal epithelial cells. *Gut* 2008; **57**: 374–82.
- 29 Chung B, Chaston T, Marks J, Srai SK, Sharp PA. Hepcidin Decreases Iron Transporter Expression in Vivo in Mouse Duodenum and Spleen and in Vitro in THP-1 Macrophages and Intestinal Caco-2 Cells. *J Nutr* 2009; **139**: 1457–62.
- 30 Arezes J, Jung G, Gabayan V, *et al.* Hepcidin-induced hypoferremia is a critical host defense mechanism against the siderophilic bacterium vibrio vulnificus. *Cell Host Microbe* 2015; **17**: 47–57.
- 31 Stefanova D, Raychev A, Arezes J, *et al.* Endogenous hepcidin and its agonist mediate resistance to selected infections by clearing non–transferrin-bound iron. *Blood* 2017; **130**: 245–57.
- 32 Michels KR, Zhang Z, Bettina AM, *et al.* Hepcidin-mediated iron sequestration protects against bacterial dissemination during pneumonia. *JCI insight* 2017; **2**: e92002.
- 33 Prentice S, Jallow AT, Sinjanka E, *et al.* Hepcidin mediates hypoferremia and reduces the growth potential of bacteria in the immediate post-natal period in human neonates. *Sci Rep* 2019.

- 34 van der Vorm LN, Hendriks JCM, Laarakkers CM, *et al.* Toward Worldwide Hepcidin Assay Harmonization: Identification of a Commutable Secondary Reference Material. *Clin Chem* 2016; **62**: 993–1001.
- 35 UNICEF | WHO | The World Bank Group | UN Population Division. Levels & trends in child mortality: report 2018. Estimates developed by the UN Inter-agency Group for Child Mortality Estimation. 2018 <https://childmortality.org/wp-content/uploads/2018/12/UN-IGME-Child-Mortality-Report-2018.pdf>.
- 36 Lawn JE, Blencowe H, Oza S, *et al.* Every newborn: Progress, priorities, and potential beyond survival. *Lancet*. 2014; **384**: 189–205.
- 37 Lawn JE, Cousens S, Zupan J. 4 Million neonatal deaths: When? Where? Why? *Lancet*. 2005; **365**: 891–900.
- 38 UNICEF. Data - Progress for every child in the SDG era. 2018. Report. [https://www.unicef.org/media/48066/file/Progress\\_for\\_Every\\_Child\\_in\\_the\\_SDG\\_Era.pdf](https://www.unicef.org/media/48066/file/Progress_for_Every_Child_in_the_SDG_Era.pdf).
- 39 Cousens S, Blencowe H, Stanton C, *et al.* National, regional, and worldwide estimates of stillbirth rates in 2009 with trends since 1995: A systematic analysis. *Lancet* 2011; **377**: 1319–30.
- 40 Kerber KJ, de Graft-Johnson JE, qar Bhutta ZA, Okong P, Starrs A, Lawn JE. Continuum of care for maternal, newborn, and child health: from slogan to service delivery. 2007; **13**; 370:1358-69.
- 41 WHO | Newborns: reducing mortality. 2016. Report <https://www.who.int/news-room/fact-sheets/detail/newborns-reducing-mortality>.
- 42 Peña-Rosas JP, De-Regil LM, Dowswell T, Viteri FE. Daily oral iron supplementation during pregnancy. In: Cochrane Database of Systematic Reviews. John Wiley & Sons, Ltd, 2012; **12**: ced004736.
- 43 Tunçalp Ö, Pena-Rosas J, Lawrie T, *et al.* WHO recommendations on antenatal care for a positive pregnancy experience-going beyond survival. *BJOG An Int J Obstet Gynaecol* 2017; **124**: 860–2.

- 44 UNICEF Data. Antenatal care. 2008 Report <https://data.unicef.org/topic/maternal-health/antenatal-care/> (accessed Dec 12, 2019).
- 45 Tikmani SS, Ali SA, Saleem S, *et al.* Trends of antenatal care during pregnancy in low- and middle-income countries: Findings from the global network maternal and newborn health registry. *Semin. Perinatol.* 2019; **43**: 297–307.
- 46 Downe S, Finlayson K, Tunçalp Ö, Gülmezoglu AM. Provision and uptake of routine antenatal services: A qualitative evidence synthesis. *Cochrane Database Syst. Rev.* 2019; **6**: cd012392
- 47 WHO. Making pregnancy safer: the critical role of the skilled attendant. A joint statement by WHO, ICM and FIGO. *Geneva, Switz WHO* 2004; 18.  
<https://apps.who.int/iris/bitstream/handle/10665/42955/9241591692.pdf?sequence=1>
- 48 Zielinski R, Ackerson K, Low LK. Planned home birth: Benefits, risks, and opportunities. *Int. J. Womens. Health.* 2015; **7**: 361–77.
- 49 Montagu D, Sudhinaraset M, Diamond-Smith N, *et al.* Where women go to deliver: understanding the changing landscape of childbirth in Africa and Asia. *Health Policy Plan* 2017; **32**: 1146–52.
- 50 Cross S, Gon G, Morrison E, *et al.* Global Health Action An invisible workforce: the neglected role of cleaners in patient safety on maternity units) An invisible workforce: the neglected role of cleaners in patient safety on maternity units. *Glob Health Action* 1480; **12**: 1480085.
- 51 Essendi H, Johnson FA, Madise N, *et al.* Infrastructural challenges to better health in maternity facilities in rural Kenya: Community and healthworker perceptions. *Reprod Health* 2015; **12**: 103.
- 52 Moyimane MB, Matlala SF, Kekana MP. Experiences of nurses on the critical shortage of medical equipment at a rural district hospital in South Africa: A qualitative study. *Pan Afr Med J* 2017; **28**: 100.
- 53 Graham WJ, Morrison E, Dancer S, *et al.* What are the threats from antimicrobial resistance for maternity units in low- and middle- income countries? *Glob Health*



- Action* 2016; **9**: 33381.
- 54 Cross S, Afsana K, Banu M, *et al.* Hygiene on maternity units: lessons from a needs assessment in Bangladesh and India. *Glob Health Action* 2016; **9**: 32541.
  - 55 Gerein N, Green A, Pearson S. Reproductive Health Matters An international journal on sexual and reproductive health and rights The Implications of Shortages of Health Professionals for Maternal Health in Sub-Saharan Africa. *Matern Heal Sub-Saharan Africa* 2006; **14**: 40–50.
  - 56 Sankar MJ, Natarajan CK, Das RR, Agarwal R, Chandrasekaran A, Paul VK. When do newborns die? A systematic review of timing of overall and cause-specific neonatal deaths in developing countries. *J. Perinatol.* 2016; **36**: S1–11.
  - 57 Seidman G, Unnikrishnan S, Kenny E, *et al.* Barriers and enablers of Kangaroo mother care practice: A systematic review. *PLoS One* 2015; **10**: e0125643.
  - 58 Mallick L, Yourkavitch J, Allen C. Trends, determinants, and newborn mortality related to thermal care and umbilical cord care practices in South Asia. *BMC Pediatr* 2019; **19**: 248.
  - 59 Khatun H, Comins CA, Shah R, Munirul Islam M, Choudhury N, Ahmed T. Uncovering the barriers to exclusive breastfeeding for mothers living in Dhaka's slums: a mixed method study. *Int Breastfeed J* 2018; **13**: 44.
  - 60 Gelano TF, Bacha YD, Abate D. Effect of chlorhexidine cord application on prevention of neonatal sepsis in developing countries: Systematic review and meta-analysis. *Int J Health Sci (Qassim)*; **13**: 40–51.
  - 61 Conde-Agudelo A, Díaz-Rossello JL. Kangaroo mother care to reduce morbidity and mortality in low birthweight infants. *Cochrane Database Syst. Rev.* 2016; **3**: cd002771.
  - 62 Quigley MA, Carson C, Sacker A, Kelly Y. Exclusive breastfeeding duration and infant infection. *Eur J Clin Nutr* 2016; **70**: 1420–7.
  - 63 Lawn JE, Osrin D, Adler A, Cousens S. Four million neonatal deaths: Counting and attribution of cause of death. *Paediatr Perinat Epidemiol* 2008; **22**: 410–6.
  - 64 UNICEF. Every child's birth right: inequities and trends in birth registration. 2013

- <http://www.cmsny.org/wp-content/uploads/Cappa.pdf> (accessed Dec 13, 2019).
- 65 Akseer N, Lawn JE, Keenan W, *et al.* Ending preventable newborn deaths in a generation. *Int J Gynaecol Obstet* 2015; : S43-8.
  - 66 United Nations. The Millennium Development Goals Report. 2015.  
[https://www.un.org/millenniumgoals/2015\\_MDG\\_Report/pdf/MDG%202015%20rev%20\(July%201\).pdf](https://www.un.org/millenniumgoals/2015_MDG_Report/pdf/MDG%202015%20rev%20(July%201).pdf)
  - 67 Countdown to 2030. Countdown to 2030 - Maternal, Newborn & Child Survival. 2018.  
<https://www.healthynewbornnetwork.org/hnn-content/uploads/Gambia-CD2030.pdf> (accessed Dec 12, 2019).
  - 68 Healthy Newborn Network. The Gambia.  
<https://www.healthynewbornnetwork.org/country/gambia/> (accessed Dec 12, 2019).
  - 69 WHO. The Gambia | Country Cooperation Strategy - at a glance. 2018.  
<https://www.afro.who.int/publications/country-cooperation-strategy-glance-gambia>
  - 70 Gambia Bureau of Statistics. The Gambia Demographic and Health Survey 2013  
<https://dhsprogram.com/pubs/pdf/FR289/FR289.pdf> (accessed Dec 9, 2019).
  - 71 Ndidi E, Oseremen I. Reasons given by pregnant women for late initiation of antenatal care in the Niger Delta, Nigeria. *Ghana Med J* 2011; **44**: 47-51
  - 72 Anya SE, Hydera A, Jaiteh LE. Antenatal care in The Gambia: Missed opportunity for information, education and communication. *BMC Pregnancy Childbirth* 2008; **8**: 9.
  - 73 Bondevik GT, Lie RT, Ulstein M, Kvale G. Maternal hematological status and risk of low birth weight and preterm delivery in Nepal. *Acta Obstet Gynecol Scand* 2001; **80**: 402–8.
  - 74 Lawn J, Mongi P, Cousens S. Africa's newborns – counting them and making them count. *WHO Report - Opportunities for Africa's Newborns* 2006.  
<https://www.who.int/pmnch/media/publications/oanfullreport.pdf>
  - 75 Ranjeva SL, Warf BC, Schiff SJ. Economic burden of neonatal sepsis in sub-Saharan Africa. *BMJ Glob Heal* 2018; **3**: e000347.
  - 76 Mwaniki MK, Atieno M, Lawn JE, Newton CRJC. Long-term neurodevelopmental

- outcomes after intrauterine and neonatal insults: A systematic review. *Lancet* 2012; **379**: 445–52.
- 77 Simonsen KA, Anderson-Berry AL, Delair SF, Davies HD. Early-onset neonatal sepsis. *Clin Microbiol Rev* 2014; **27**: 21–47.
  - 78 Dong Y, Speer CP. Late-onset neonatal sepsis:Recent developments. *Arch. Dis. Child. Fetal Neonatal Ed.* 2015; **100**: F257–63.
  - 79 Okomo U, Akpalu NK, Le Doare K, *et al.* Aetiology of invasive bacterial infection and antimicrobial resistance in neonates in sub-Saharan Africa: a systematic review and meta-analysis in line with the STROBE-NI reporting guidelines. *Lancet Infect Dis.* 2019; **11**: 1219-1234
  - 80 Mulholland EK, Ogunlesi OO, Adegbola RA, *et al.* Etiology of serious infections in young Gambian infants. *Pediatr Infect Dis J* 1999; **18**: S35-41.
  - 81 Downey LC, Smith PB, Benjamin DK. Risk factors and prevention of late-onset sepsis in premature infants. *Early Hum Dev* 2010; **86**: 7–12.
  - 82 Jimenez-Truque N, Tedeschi S, Saye EJ, *et al.* Relationship between maternal and neonatal *Staphylococcus aureus* colonization. *Pediatrics* 2012; **129**: e1252-9.
  - 83 Romano-Bertrand S, Filleron A, Mesnage R, *et al.* *Staphylococcus aureus* in a neonatal care center: methicillin-susceptible strains should be a main concern. *Antimicrob Resist Infect Control* 2014; **3**: 21.
  - 84 Hamer DH, Darmstadt GL, Carlin JB, *et al.* Etiology of Bacteremia in Young Infants in Six Countries. *Pediatr Infect Dis J* 2015; **34**: e1–8.
  - 85 van Hal SJ, Jensen SO, Vaska VL, Espedido BA, Paterson DL, Gosbell IB. Predictors of Mortality in *Staphylococcus aureus* Bacteremia. *Clin Microbiol Rev* 2012; **25**: 362–86.
  - 86 Nelson MU, Gallagher PG. Methicillin-Resistant *Staphylococcus aureus* in the Neonatal Intensive Care Unit. *Semin. Perinatol.* 2012; **36**: 424–30.
  - 87 Denkel LA, Schwab F, Kola A, *et al.* The mother as most important risk factor for colonization of very low birth weight (VLBW) infants with extended-spectrum b-

- lactamase-producing Enterobacteriaceae (ESBL-E). *J Antimicrob Chemother* 2014; **69**: 2230–7.
- 88 Podschun R, Ullmann U. Klebsiella spp. as nosocomial pathogens: epidemiology, taxonomy, typing methods, and pathogenicity factors. *Clin Microbiol Rev* 1998; **11**: 589–603.
  - 89 Seifi K, Kazemian H, Heidari H, *et al.* Evaluation of biofilm formation among klebsiella pneumoniae isolates and molecular characterization by ERIC-PCR. *Jundishapur J Microbiol* 2016; **9**: e30682.
  - 90 Dsouza R, Pinto NA, Hwang IS, *et al.* Panel strain of Klebsiella pneumoniae for beta-lactam antibiotic evaluation: Their phenotypic and genotypic characterization. *PeerJ* 2017; **5**: e2896.
  - 91 Peigne C, Bidet P, Mahjoub-Messai F, *et al.* The plasmid of Escherichia coli strain S88 (O45:K1:H7) that causes neonatal meningitis is closely related to avian pathogenic E. coli plasmids and is associated with high-level bacteremia in a neonatal rat meningitis model. *Infect Immun* 2009; **77**: 2272–84.
  - 92 Zea-Vera A, Ochoa TJ. Challenges in the diagnosis and management of neonatal sepsis. *J Trop Pediatr* 2015; **61**: 1–13.
  - 93 Saha SK, Schrag SJ, El Arifeen S, *et al.* Causes and incidence of community-acquired serious infections among young children in south Asia (ANISA): an observational cohort study. *Lancet (London, England)* 2018; **392**: 145–59.
  - 94 Venkatesh M, Flores A, Luna RA, Versalovic J. Molecular microbiological methods in the diagnosis of neonatal sepsis. *Expert Rev Anti Infect Ther* 2010; **8**: 1037–48.
  - 95 Young Infants Clinical Signs Study Group. Clinical signs that predict severe illness in children under age 2 months: a multicentre study. *Lancet* 2008; **371**: 135–42.
  - 96 Seale AC, Blencowe H, Manu AA, *et al.* Estimates of possible severe bacterial infection in neonates in sub-Saharan Africa, south Asia, and Latin America for 2012: A systematic review and meta-analysis. *Lancet Infect Dis* 2014; **14**: 731–41.
  - 97 Simonsen KA, Anderson-Berry AL, Delair SF, Davies HD. Early-Onset Neonatal

- Sepsis. *Clin Microbiol Rev* 2014; **27**: 21–47.
- 98 Fuchs A, Bielicki J, Mathur S, Sharland M, Van Den Anker JN. Reviewing the WHO guidelines for antibiotic use for sepsis in neonates and children. *Paediatr. Int. Child Health*. 2018; **38**: S3–15.
  - 99 Darmstadt GL, Batra M, Zaidi AKM. Oral Antibiotics in the Management of Serious Neonatal Bacterial Infections in Developing Country Communities. *Pediatr Infect Dis J* 2009; **28**: S31–6.
  - 100 O'Neill J. Antimicrobial Resistance : Tackling a crisis for the health and wealth of nations. *Rev Antimicrob Resist* 2014; 1–16.
  - 101 Downie L, Armiento R, Subhi R, Kelly J, Clifford V, Duke T. Community-acquired neonatal and infant sepsis in developing countries: Efficacy of WHO's currently recommended antibiotics - Systematic review and meta-analysis. *Arch Dis Child* 2013; **98**: 146–54.
  - 102 Laxminarayan R, Matsoso P, Pant S, *et al*. Access to effective antimicrobials: A worldwide challenge. *Lancet* 2016; **387**: 168–75.
  - 103 Kayange N, Kamugisha E, Mwizamholya DL, Jeremiah S, Mshana SE. Predictors of positive blood culture and deaths among neonates with suspected neonatal sepsis in a tertiary hospital, Mwanza- Tanzania. *BMC Pediatr* 2010; **10**: 39
  - 104 Klingenberg C, Olomi R, Oneko M, Sam N, Langeland N. Neonatal morbidity and mortality in a Tanzanian tertiary care referral hospital. *Ann Trop Paediatr* 2003; **23**: 293–9.
  - 105 Exner M, Bhattacharya S, Christiansen B, *et al*. Antibiotic resistance: What is so special about multidrug-resistant Gram-negative bacteria? *GMS Hyg Infect Control* 2017; **12**: Doc05.
  - 106 Iredell J, Brown J, Tagg K. Antibiotic resistance in Enterobacteriaceae: Mechanisms and clinical implications. *BMJ*. 2016; **352**: h6420
  - 107 Laxminarayan R, Heymann DL. Challenges of drug resistance in the developing world. *BMJ*. 2012; **344**: e1567.

- 108 Morrill HJ, Pogue JM, Kaye KS, LaPlante KL. Treatment options for carbapenem-resistant Enterobacteriaceae infections. *Open Forum Infect Dis* 2015; **2**: ofv050.
- 109 Bonomo RA. New Delhi metallo-B-lactamase and multidrug resistance: A global SOS? *Clin Infect Dis* 2011; **52**: 485–7.
- 110 Arnold RS, Thom KA, Sharma S, Phillips M, Kristie Johnson J, Morgan DJ. Emergence of *Klebsiella pneumoniae* carbapenemase-producing bacteria. *South Med J* 2011; **104**: 40–5.
- 111 Datta S, Roy S, Chatterjee S, *et al.* A Five-Year Experience of Carbapenem Resistance in Enterobacteriaceae Causing Neonatal Septicaemia: Predominance of NDM-1. *PLoS One* 2014; **9**: e112101.
- 112 Ballot DE, Bandini R, Nana T, *et al.* A review of -multidrug-resistant Enterobacteriaceae in a neonatal unit in Johannesburg, South Africa. *BMC Pediatr* 2019; **19**: 320.
- 113 Chambers HF, DeLeo FR. Waves of resistance: *Staphylococcus aureus* in the antibiotic era. *Nat. Rev. Microbiol.* 2009; **7**: 629–41.
- 114 Uhlemann A-C, Otto M, Lowy FD, DeLeo FR. Evolution of community- and healthcare-associated methicillin-resistant *Staphylococcus aureus*. *Infect Genet Evol* 2014; **21**: 563–74.
- 115 David MZ, Daum RS. Community-associated methicillin-resistant *Staphylococcus aureus*: Epidemiology and clinical consequences of an emerging epidemic. *Clin. Microbiol. Rev.* 2010; **23**: 616–87.
- 116 Mwananyanda L, Pierre C, Mwansa J, *et al.* Preventing Bloodstream Infections and Death in Zambian Neonates: Impact of a Low-cost Infection Control Bundle. *Clin Infect Dis* 2019; **69**: 1360–7.
- 117 Roca A, Oluwalana C, Bojang A, *et al.* Oral azithromycin given during labour decreases bacterial carriage in the mothers and their offspring: a double-blind randomized trial. *Clin Microbiol Infect* 2016; **22**: 565.e1-565.e9.
- 118 Unger HW, Ome-Kaius M, Wangnapi RA, *et al.* Sulphadoxine-pyrimethamine plus

- azithromycin for the prevention of low birthweight in Papua New Guinea: a randomised controlled trial. *BMC Med* 2015; **13**: 9.
- 119 Keenan JD, Bailey RL, West SK, *et al.* Azithromycin to Reduce Childhood Mortality in Sub-Saharan Africa. *N Engl J Med* 2018; **378**: 1583–92.
  - 120 Bojang A, Camara B, Jagne Cox I, *et al.* Long-term Impact of Oral Azithromycin Taken by Gambian Women During Labor on Prevalence and Antibiotic Susceptibility of *Streptococcus pneumoniae* and *Staphylococcus aureus* in Their Infants: Follow-up of a Randomized Clinical Trial. *Clin Infect Dis* 2018; **67**: 1191–7.
  - 121 Renz H, Brandtzaeg P, Hornef M. The impact of perinatal immune development on mucosal homeostasis and chronic inflammation. *Nat. Rev. Immunol.* 2012; **12**: 9–23.
  - 122 Al Faleh K, Anabrees J. Probiotics for prevention of necrotizing enterocolitis in preterm infants. *Cochrane Database Syst Rev* 2014; cd005496
  - 123 Nandhini LP, Biswal N, Adhisivam B, Mandal J, Bhatb V, Mathai B. Synbiotics for decreasing incidence of necrotizing enterocolitis among preterm neonates - A randomized controlled trial. *J Matern Neonatal Med* 2016; **29**: 821–5.
  - 124 McElhinney DB, Hedrick HL, Bush DM, *et al.* Necrotizing enterocolitis in neonates with congenital heart disease: Risk factors and outcomes. *Pediatrics* 2000; **106**: 1080–7.
  - 125 Dilli D, Aydin B, Fettah ND, *et al.* The pro-pre-save study: Effects of probiotics and prebiotics alone or combined on necrotizing enterocolitis in very low birth weight infants. *J Pediatr* 2015; **166**: 545-551.e1.
  - 126 Panigrahi P, Parida S, Nanda NC, *et al.* A randomized synbiotic trial to prevent sepsis among infants in rural India. *Nature* 2017; **548**: 407–12.
  - 127 Archibald F. *Lactobacillus plantarum*, an organism not requiring iron. *FEMS Microbiol Lett* 1983; **19**: 29–32.
  - 128 Hillman NH, Kallapur SG, Jobe AH. Physiology of transition from intrauterine to extrauterine life. *Clin. Perinatol.* 2012; **39**: 769–83.
  - 129 Stinson LF, Boyce MC, Payne MS, Keelan JA. The not-so-sterile womb: Evidence

- that the human fetus is exposed to bacteria prior to birth. *Front Microbiol* 2019; **10**: 1124.
- 130 Perez-Muñoz ME, Arrieta MC, Ramer-Tait AE, Walter J. A critical assessment of the 'sterile womb' and 'in utero colonization' hypotheses: Implications for research on the pioneer infant microbiome. *Microbiome*. 2017; **5**: 48
  - 131 Kates EH, Kates JS. Anemia and polycythemia in the newborn. *Pediatr. Rev.* 2007; **28**: 33–4.
  - 132 Szabo M, Vasarhelyi B, Balla G, *et al.* Acute postnatal increase of extracellular antioxidant defence of neonates: the role of iron metabolism. *Acta Paediatr* 2001; **90**: 1167–70.
  - 133 Sankaran VG, Xu J, Orkin SH. Advances in the understanding of haemoglobin switching. *Br J Haematol* 2010; **149**: 181–94.
  - 134 Chou SC, Palmer RH, Ezhuthachan S, *et al.* Management of Hyperbilirubinemia in Newborns: Measuring Performance by Using a Benchmarking Model. *Pediatrics* 2003; **112**: 1264–73.
  - 135 Mogensen TH. Pathogen recognition and inflammatory signaling in innate immune defenses. *Clin. Microbiol. Rev.* 2009; **22**: 240–73.
  - 136 Basha S, Surendran N, Pichichero M. Immune responses in neonates. *Expert Rev Clin Immunol* 2014; **10**: 1171–84.
  - 137 Yost CC, Cody MJ, Harris ES, *et al.* Impaired neutrophil extracellular trap (NET) formation: a novel innate immune deficiency of human neonates. *Blood* 2009; **113**: 6419–27.
  - 138 Carr R. Neutrophil production and function in newborn infants. *Br. J. Haematol.* 2000; **110**: 18–28.
  - 139 Singh G, Archana G. Unraveling the mystery of vernix caseosa. *Indian J. Dermatol.* 2008; **53**: 54–60.
  - 140 Melville JM, Moss TJM. The immune consequences of preterm birth. *Front. Neurosci.* 2013; **7**: 79.



- 141 Fietta A, Sacchi F, Bersani C, *et al.* Complement-dependent bactericidal activity for *E. coli* K12 in serum of preterm newborn infants. *Acta Paediatr Scand* 1987; **76**: 37–41.
- 142 Proytcheva MA. Issues in Neonatal Cellular Analysis. *Am J Clin Pathol* 2009; **131**: 560–73.
- 143 Hood MI, Skaar EP. Nutritional immunity: transition metals at the pathogen-host interface. *Nat Rev Microbiol* 2012; **10**: 525–37.
- 144 Cherayil BJ. Iron and immunity: Immunological consequences of iron deficiency and overload. *Arch. Immunol. Ther. Exp. (Warsz)*. 2010; **58**: 407–15.
- 145 Cassat JE, Skaar EP. Iron in infection and immunity. *Cell Host Microbe*. 2013; **13**: 509–19.
- 146 Caza M, Kronstad JW. Shared and distinct mechanisms of iron acquisition by bacterial and fungal pathogens of humans. *Front Cell Infect Microbiol* 2013; **3**: 80.
- 147 Koczura R, Kaznowski A. Occurrence of the *Yersinia* high-pathogenicity island and iron uptake systems in clinical isolates of *Klebsiella pneumoniae*. *Microb Pathog* 2003; **35**: 197–202.
- 148 Muryoi N, Tiedemann MT, Pluym M, Cheung J, Heinrichs DE, Stillman MJ. Demonstration of the iron-regulated surface determinant (Isd) heme transfer pathway in *Staphylococcus aureus*. *J Biol Chem* 2008; **283**: 28125–36.
- 149 Grigg JC, Ukpabi G, Gaudin CFM, Murphy MEP. Structural biology of heme binding in the *Staphylococcus aureus* Isd system. *J Inorg Biochem* 2010; **104**: 341–8.
- 150 Maresso AW, Schneewind O. Iron acquisition and transport in *Staphylococcus aureus*. *Biometals* 2006; **19**: 193–203.
- 151 Law D, Kelly J. Use of heme and hemoglobin by *Escherichia coli* O157 and other Shiga-like-toxin-producing *E. coli* serogroups. *Infect Immun* 1995; **63**: 700–2.
- 152 Runyen-Janecky LJ. Role and regulation of heme iron acquisition in gram-negative pathogens. *Front Cell Infect Microbiol* 2013; **3**: 55.
- 153 Hagan EC, Mobley HLT. Haem acquisition is facilitated by a novel receptor Hma and required by uropathogenic *Escherichia coli* for kidney infection. *Mol Microbiol* 2009;

- 71: 79–91.
- 154 Romero-Espejel ME, González-López MA, Olivares-Trejo J de J. Streptococcus pneumoniae requires iron for its viability and expresses two membrane proteins that bind haemoglobin and haem. *Metallomics* 2013; **5**: 384–9.
- 155 Tai SS, Lee CJ, Winter RE. Hemin utilization is related to virulence of Streptococcus pneumoniae. *Infect Immun* 1993; **61**: 5401–5.
- 156 Vázquez-Zamorano ZE, González-López MA, Romero-Espejel ME, Azuara-Liceaga EI, López-Casamichana M, Olivares-Trejo J de J. Streptococcus pneumoniae secretes a glyceraldehyde-3-phosphate dehydrogenase, which binds haemoglobin and haem. *Biometals* 2014; **27**: 683–93.
- 157 Fernandez A, Lechardeur D, Derré-Bobillot A, Couvé E, Gaudu P, Gruss A. Two coregulated efflux transporters modulate intracellular heme and protoporphyrin IX availability in Streptococcus agalactiae. *PLoS Pathog* 2010; **6**: e1000860.
- 158 Pishchany G, Sheldon JR, Dickson CF, *et al.* IsdB-dependent hemoglobin binding is required for acquisition of heme by Staphylococcus aureus. *J Infect Dis* 2014; **209**: 1764–72.
- 159 Torres VJ, Pishchany G, Humayun M, Schneewind O, Skaar EP. Staphylococcus aureus IsdB is a hemoglobin receptor required for heme iron utilization. *J Bacteriol* 2006; **188**: 8421–9.
- 160 Farrand AJ, Reniere ML, Ingmer H, Frees D, Skaar EP. Regulation of host hemoglobin binding by the Staphylococcus aureus Clp proteolytic system. *J Bacteriol* 2013; **195**: 5041–50.
- 161 Otto BR, van Dooren SJ, Nuijens JH, Luirink J, Oudega B. Characterization of a hemoglobin protease secreted by the pathogenic Escherichia coli strain EB1. *J Exp Med* 1998; **188**: 1091–103.
- 162 Dryla A, Hoffmann B, Gelbmann D, *et al.* High-affinity binding of the staphylococcal HarA protein to haptoglobin and hemoglobin involves a domain with an antiparallel eight-stranded ??-barrel fold. *J Bacteriol* 2007; **189**: 254–64.

- 163 Dryla A, Gelbmann D, von Gabain A, Nagy E. Identification of a novel iron regulated staphylococcal surface protein with haptoglobin-haemoglobin binding activity. *Mol Microbiol* 2003; **49**: 37–53.
- 164 Francis RT, Booth JW, Becker RR. Uptake of iron from hemoglobin and the haptoglobin-hemoglobin complex by hemolytic bacteria. *Int J Biochem* 1985; **17**: 767–73.
- 165 Beasley FC. Serum iron uptake and virulence in *Staphylococcus aureus*. *Electron Thesis Diss Repos* 2011. <http://ir.lib.uwo.ca/etd/135> (accessed Sept 12, 2016).
- 166 Park R-Y, Sun H-Y, Choi M-H, Bai Y-H, Shin S-H. *Staphylococcus aureus* siderophore-mediated iron-acquisition system plays a dominant and essential role in the utilization of transferrin-bound iron. *J Microbiol* 2005; **43**: 183–90.
- 167 Freestone PPE, Haigh RD, Williams PH, Lyte M. Involvement of enterobactin in norepinephrine-mediated iron supply from transferrin to enterohaemorrhagic *Escherichia coli*. *FEMS Microbiol Lett* 2003; **222**: 39–43.
- 168 Caza M, Lépine F, Milot S, Dozois CM. Specific roles of the *iroBCDEN* genes in virulence of an avian pathogenic *Escherichia coli* O78 strain and in production of salmochelins. *Infect Immun* 2008; **76**: 3539–49.
- 169 de Lorenzo V, Bindereif A, Paw BH, Neilands JB. Aerobactin biosynthesis and transport genes of plasmid ColV-K30 in *Escherichia coli* K-12. *J Bacteriol* 1986; **165**: 570–8.
- 170 Cornelissen CN, Sparling PF. Iron piracy: acquisition of transferrin-bound iron by bacterial pathogens. *Mol Microbiol* 1994; **14**: 843–50.
- 171 Lawlor MS, O'Connor C, Miller VL. Yersiniabactin is a virulence factor for *Klebsiella pneumoniae* during pulmonary infection. *Infect Immun* 2007; **75**: 1463–72.
- 172 Bachman MA, Miller VL, Weiser JN. Mucosal lipocalin 2 has pro-inflammatory and iron-sequestering effects in response to bacterial enterobactin. *PLoS Pathog* 2009; **5**: e1000622.
- 173 Perry RD, Clemente CLS. Siderophore synthesis in *Klebsiella pneumoniae* and

- Shigella sonnei during iron Siderophore Synthesis in Klebsiella pneumoniae and Shigella sonnei During Iron Deficiency. *J Bacteriol* 1979; **140**: 1129–32.
- 174 Hsieh P-F, Lin T-L, Lee C-Z, Tsai S-F, Wang J-T. Serum-induced iron-acquisition systems and TonB contribute to virulence in Klebsiella pneumoniae causing primary pyogenic liver abscess. *J Infect Dis* 2008; **197**: 1717–27.
- 175 Clancy A, Loar JW, Speziali CD, Oberg M, Heinrichs DE, Rubens CE. Evidence for siderophore-dependent iron acquisition in group B streptococcus. *Mol Microbiol* 2006; **59**: 707–21.
- 176 Modun B, Morrissey J, Williams P. The staphylococcal transferrin receptor: A glycolytic enzyme with novel functions. *Trends Microbiol.* 2000; **8**: 231–7.
- 177 Taylor JM, Heinrichs DE. Transferrin binding in Staphylococcus aureus: involvement of a cell wall-anchored protein. *Mol Microbiol* 2002; **43**: 1603–14.
- 178 Sandrini S, Masania R, Zia F, Haigh R, Freestone P. Role of porin proteins in acquisition of transferrin iron by enteropathogens. *Microbiol* 2013; **159**: 2639–50.
- 179 Lisiecki P, Mikucki J. Human body iron sources utilized in vitro by staphylococci. *Med doświadczalna i Mikrobiol* 1996; **48**: 5–13.
- 180 Clarke SR, Foster SJ. IsdA protects Staphylococcus aureus against the bactericidal protease activity of apolactoferrin. *Infect Immun* 2008; **76**: 1518–26.
- 181 Konetschny-Rapp S, Jung G, Meiwes J, Zahner H. Staphyloferrin A: A structurally new siderophore from staphylococci. *Eur J Biochem* 1990; **191**: 65–74.
- 182 Braun V, Braun M. Iron transport and signaling in Escherichia coli. *FEBS Lett* 2002; **529**: 78–85.
- 183 Brown JS, Gilliland SM, Holden DW. A Streptococcus pneumoniae pathogenicity island encoding an ABC transporter involved in iron uptake and virulence. *Mol Microbiol* 2001; **40**: 572–85.
- 184 Yang X-Y, He K, Du G, *et al.* Integrated Translatomics with Proteomics to Identify Novel Iron-Transporting Proteins in Streptococcus pneumoniae. *Front Microbiol* 2016; **7**: 78.

- 185 Horsburgh MJ, Ingham E, Foster SJ. In *Staphylococcus aureus*, fur is an interactive regulator with PerR, contributes to virulence, and is necessary for oxidative stress resistance through positive regulation of catalase and iron homeostasis. *J Bacteriol* 2001; **183**: 468–75.
- 186 Lau CKY, Ishida H, Liu Z, Vogel HJ. Solution structure of *Escherichia coli* FeoA and its potential role in bacterial ferrous iron transport. *J Bacteriol* 2013; **195**: 46–55.
- 187 Kammler M, Schön C, Hantke K. Characterization of the ferrous iron uptake system of *Escherichia coli*. *J Bacteriol* 1993; **175**: 6212–9.
- 188 Grosse C, Scherer J, Koch D, Otto M, Taudte N, Grass G. A new ferrous iron-uptake transporter, EfeU (YcdN), from *Escherichia coli*. *Mol Microbiol* 2006; **62**: 120–31.
- 189 Hung K-W, Juan T-H, Hsu Y-L, Huang TH. NMR structure note: the ferrous iron transport protein C (FeoC) from *Klebsiella pneumoniae*. *J Biomol NMR* 2012; **53**: 161–5.
- 190 Li-Ching Kok. Comparative characterization of the ferrous iron transport systems FeoABC, SitABCD and EfeUOB in *Klebsiella pneumoniae* CG43. 2014.  
<https://ir.nctu.edu.tw/bitstream/11536/75800/1/703301.pdf>
- 191 Bray BA, Sutcliffe IC, Harrington DJ. Expression of the MtsA lipoprotein of *Streptococcus agalactiae* A909 is regulated by manganese and iron. *Int J Gen Mol Microbiol* 2009; **95**: 101–9.
- 192 Tidmarsh GF, Klebba PE, Rosenberg LT. Rapid release of iron from ferritin by siderophores. *J Inorg Biochem* 1983; **18**: 161–8.
- 193 Barton Pai A, Pai MP, Depczynski J, McQuade CR, Mercier R-C. Non-transferrin-bound iron is associated with enhanced *Staphylococcus aureus* growth in hemodialysis patients receiving intravenous iron sucrose. *Am J Nephrol* 2006; **26**: 304–9.
- 194 Brock JH, Williams PH, Licéaga J, Wooldridge KG. Relative availability of transferrin-bound iron and cell-derived iron to aerobactin-producing and enterochelin-producing strains of *Escherichia coli* and to other microorganisms. *Infect Immun* 1991; **59**: 3185–

- 90.
- 195 Kim C, Park R, Choi M, Sun H, Shin S. Ferrophilic characteristics of *Vibrio vulnificus* and potential usefulness of iron chelation therapy. *J Infect Dis* 2007; **195**: 90–8.
- 196 Cornelis P, Andrews SC. Iron uptake and homeostasis in microorganisms. Caister Academic Press, 2010.
- 197 Krause A, Neitz S, Mägert HJ, *et al.* LEAP-1, a novel highly disulfide-bonded human peptide, exhibits antimicrobial activity. *FEBS Lett* 2000; **480**: 147–50.
- 198 Pigeon C, Ilyin G, Courselaud B, *et al.* A new mouse liver-specific gene, encoding a protein homologous to human antimicrobial peptide hepcidin, is overexpressed during iron overload. *J Biol Chem* 2001; **276**: 7811–9.
- 199 Park CH, Valore E V., Waring AJ, Ganz T. Hepcidin, a Urinary Antimicrobial Peptide Synthesized in the Liver. *J Biol Chem* 2001; **276**: 7806–10.
- 200 Ganz T, Nemeth E. Iron homeostasis in host defence and inflammation. *Nat Rev Immunol* 2015; **15**: 500–10.
- 201 Ganz T, Nemeth E. Iron Sequestration and Anemia of Inflammation. *Semin Hematol* 2009; **46**: 387–93.
- 202 Parrow NL, Fleming RE, Minnick MF. Sequestration and Scavenging of Iron in Infection. *Infect Immun* 2013; **81**: 3503–14.
- 203 Sazawal S, Black RE, Ramsan M, *et al.* Effects of routine prophylactic supplementation with iron and folic acid on admission to hospital and mortality in preschool children in a high malaria transmission setting: Community-based, randomised, placebo-controlled trial. *Lancet* 2006; **367**: 133–43.
- 204 Cross JH, Bradbury RS, Fulford AJ, *et al.* Oral iron acutely elevates bacterial growth in human serum. *Sci Rep* 2015; **5**: 16670.
- 205 Barry DM, Reeve AW. Increased incidence of gram-negative neonatal sepsis with intramuscular iron administration. *Pediatrics* 1977; **60**: 908–12.
- 206 Drakesmith H, Prentice AM. Hepcidin and the Iron-Infection Axis. *Science* 2012; **338**: 768–72.

- 207 Ganz T, Nemeth E. Iron Balance and the Role of Hepcidin in Chronic Kidney Disease. *Semin. Nephrol.* 2016; **36**: 87-93
- 208 Liu J, Chen S, Ye X. The effect of red blood cell transfusion on plasma hepcidin and growth differentiation factor 15 in gastric cancer patients: a prospective study. *Ann Transl Med* 2019; **7**: 43.
- 209 Harrison-Findik DD. Is the iron regulatory hormone hepcidin a risk factor for alcoholic liver disease? *World J Gastroenterol* 2009; **15**: 1186–93.
- 210 Girelli D, Pasino M, Goodnough JB, *et al.* Reduced serum hepcidin levels in patients with chronic hepatitis C. *J Hepatol* 2009; **51**: 845–52.
- 211 Ilkovska B, Kotevska B, Trifunov G, Kanazirev B. Serum hepcidin reference range, gender differences, menopausal dependence and biochemical correlates in healthy subjects. *J IMAB.* 2016; **22**: 1127-1131.
- 212 Itkonen O, Parkkinen J, Stenman UH, Hämäläinen E. Preanalytical factors and reference intervals for serum hepcidin LC-MS/MS method. *Clin Chim Acta* 2012. 413: 696-701
- 213 Troutt JS, Rudling M, Persson L, *et al.* Circulating Human hepcidin-25 concentrations display a diurnal rhythm, increase with prolonged fasting, and are reduced by growth hormone administration. *Clin Chem* 2012; **58**: 1225-31
- 214 Kroot JJC, Hendriks JCM, Laarakkers CMM, *et al.* (Pre)analytical imprecision, between-subject variability, and daily variations in serum and urine hepcidin: Implications for clinical studies. *Anal Biochem* 2009; **389**: 124–9.
- 215 Konz T, Montes-Bayón M, Vaulont S. Hepcidin quantification: Methods and utility in diagnosis. *Metallomics.* 2014; **6**: 1583–90.
- 216 Piperno A, Mariani R, Trombini P, Girelli D. Hepcidin modulation in human diseases: From research to clinic. *World J. Gastroenterol.* 2009; **15**: 538-51
- 217 Ganz T, Nemeth E. Iron imports. IV. Hepcidin and regulation of body iron metabolism. *Am J Physiol - Gastrointest Liver Physiol* 2006; **290**: g199-203
- 218 Hunter HN, Bruce Fulton D, Ganz T, Vogel HJ. The solution structure of human

- hepcidin, a peptide hormone with antimicrobial activity that is involved in iron uptake and hereditary hemochromatosis. *J Biol Chem* 2002; **277**: 37597-603
- 219 Ford BA, Eby CS, Scott MG, Coyne DW. Intra-individual variability in serum hepcidin precludes its use as a marker of iron status in hemodialysis patients. *Kidney Int* 2010; **78**: 769–73.
- 220 Kemna EHJM, Tjalsma H, Podust VN, Swinkels DW. Mass spectrometry-based hepcidin measurements in serum and urine: Analytical aspects and clinical implications. *Clin Chem* 2007; **53**: 620–8.
- 221 Girelli D, Nemeth E, Swinkels DW. Hepcidin in the diagnosis of iron disorders. *Blood* 2016; **127**: 2809–13.
- 222 Diepeveen LE, Laarakkers CMM, Martos G, *et al.* Provisional standardization of hepcidin assays: creating a traceability chain with a primary reference material, candidate reference method and a commutable secondary reference material. *Clin Chem Lab Med* 2019; **57**: 864–72.



## Chapter 2 – PhD Study Setting

---

### Summary of Chapter

This chapter provides a brief geographical and economic overview of the Kanifing region and The Gambia as a whole. This is then followed by a description of the work environment at the study hospital, Kanifing General Hospital. Lastly, I describe the environment in which community visits take place around the Kanifing and Kombo regions, in the homes of study participants. All ethical, political and logistical issues faced in each of the study settings are detailed.

## 2.1 PhD Study Setting

### 2.1.1 The Republic of The Gambia

The Republic of The Gambia is the smallest mainland African nation, formed around the River Gambia. As part of West Africa, it is surrounded by Senegal and has a population of 2.3 million people (2018).<sup>1</sup> The population is growing at a rate of approximately 2.9% per year (2018).<sup>2</sup> Islam is practised by 95% of the country's population, with twelve main ethnic groups, the largest of which are Mandinka and Fula.<sup>3</sup> The Gambia has a market-based economy, focused around tourism and traditional subsistence agriculture.<sup>3</sup>



**Figure 2.1** Map of The Gambia. Arrow shows study site location at Kanifing General Hospital. *Source:* © OpenStreetMap contributors.

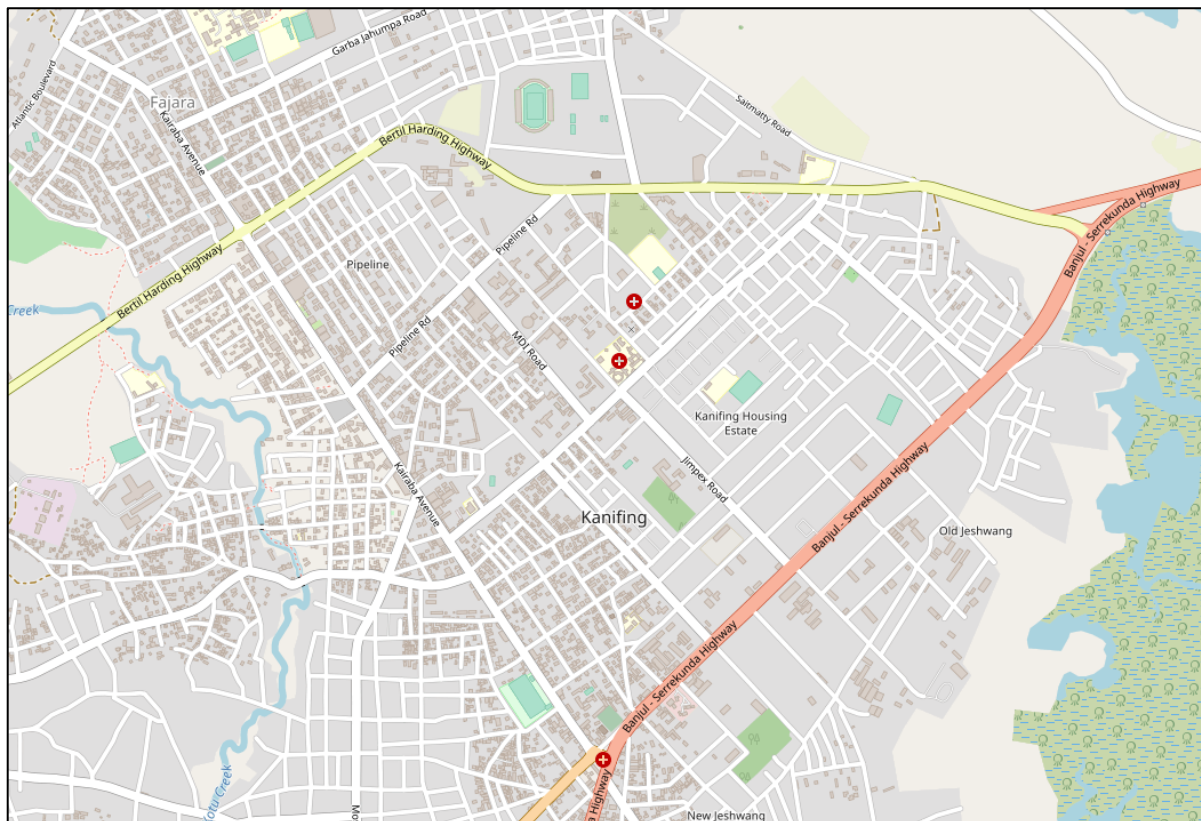
A countrywide political problem faced by the study was the removal of the previous President of The Gambia, Yahya Abdul-Aziz Jemus Junkung Jammeh. The Gambia experienced a contested election in December 2016, which provoked a constitutional crisis. The situation was resolved by military intervention from the neighbouring countries from the Economic Community of West African States (ECOWAS). The new Gambian president was inaugurated towards the end of January 2017, but even after the inauguration, the Gambian government and society were in a state of flux and political uncertainty. This affected the

timing of three aspects of the study, including ethical approvals, community sensitisation, and study initiation.

### 2.1.2 Kanifing Municipality

The Kanifing region is made up predominately of the large town, Serrekunda. Kanifing region is a peri-urban area with a population of around 370,000<sup>4</sup> and is 10 km to the southwest of the capital, Banjul. Serrekunda was initially made up of nine villages that have merged into a sprawling urban area.

### 2.1.3 Study Sites: Kanifing General Hospital, The Gambia



**Figure 2.2** Map of Kanifing, The Gambia. Source: © OpenStreetMap contributors.

Kanifing General Hospital, located in the Kanifing district, is one of six general hospitals in The Gambia. It is a public, government-run facility that first opened in 2010. Named initially Serrekunda General Hospital (SGH) it contains approximately 130 beds, with services and units such as an accident and emergency unit, out-patient unit, CT scanning department, maternity, antenatal and paediatric units. Recent investment has led to the formation of a small neonatal care unit. Complex clinical cases are commonly referred to The Edward Francis Small Teaching Hospital (EFSTH) in Banjul, as it is the only tertiary government referral hospital.<sup>3</sup> Seven paediatricians staff the maternity and paediatric wards, along with sixteen senior nurses/midwives. During the study period, usually, there were four to six trained nurses working in the maternity ward during the day shift, with an equal number of nurse attendants. Staffing levels would commonly reduce during the night shift. Previously, the hospital has experienced periodic shortages of medicines, supplies, staff, electricity and water.



**Figure 2.3** Kanifing General Hospital Maternity Ward. Neolnnate study team recruiting (*top left*), cord bleeding (*top right*), electronic data collecting (*bottom left*) at Kanifing General Hospital Maternity Ward (*bottom right*). *Note: all imaged individuals have provided photographic consent.*

Annually, Kanifing General Hospital (KGH) provides antenatal care to 500–700 pregnant mothers (unpublished - hospital data). Mothers receiving antenatal care at other local primary health-care facilities increase the total number of births at the hospital to 3500–4500 per year (unpublished - hospital data). The percentage of these that are live, low birthweight neonates (<2.5kg) is approximately 10% (hospital data 2016-2018, unpublished). During the study period, rates of stillbirth were at times poorly documented by hospital maternity ward staff, as seen in other institutions in low- and middle-income countries (LMICs).<sup>5</sup>

The maternity ward was split into two large wards, with one being used for pre- and post-delivery mothers and their neonates. There was also an adjacent eight-bed open-plan delivery ward. The delivery ward used a single partially broken neonatal resuscitator, which was commonly occupied by multiple neonates (1-3) at peak admission periods. The delivery ward also possessed limited respiratory support in the form of a single electrical oxygen concentrator. Thanks to the help from MRCG at LSHTM, both issues were overcome quickly in order to ensure service quality greatly improved. Other barriers to care included the lack of implementing Kangaroo mother care (KMC) in the hospital at the beginning of the study. As a result of a local implementation study, we were able to train several doctors and senior nurses on the care protocols. Infection control was also deemed to be poor quality in the maternity ward during the initiation of the study. As a result, we ensured new cleaning equipment was available on a monthly basis, in addition to investment into new waste systems, weighing scale, delivery equipment, storage devices and worktables.

A study-specific issue we experienced at Kanifing General Hospital was that mothers who delivered at the hospital were not always receiving antenatal care at the same health facility. At the beginning of the study, this resulted in many newborns being missed to recruitment. Anecdotal evidence suggested that mothers who were approached at antenatal visits at Kanifing General Hospital were often choosing to deliver at home or a primary level healthcare facility. Subsequently, our study team worked with local midwives and nurses to ensure mothers were informed of the benefits and risks associated with delivery at Kanifing General Hospital and being part of the Neonate Study. Furthermore, we also produced an additional recruitment route to ensure mothers receiving antenatal care outside the hospital could equally have their newborn recruited to the Neonate Study. This was a two-stage process, in which on arrival to the maternity ward if the mother was found to be capable, she would consent to the collection and storage of cord blood in the maternity ward. After a few hours, and when the mother was deemed to be in the right frame of mind after the delivery process, she was asked as to whether the cord blood sample and her neonate could be full recruited into the study. If at this point, the mother refused to give consent, the cord blood sample and her data were removed from the fridge and study server, respectively. This aspect of the study protocol required high-quality consent training focused on detailed discussions concerning different scenarios with the study staff.



#### 2.1.4 Study Sites: Community Visits (West Coast Region)



**Figure 2.4** Community study visits to the homes of recruited newborns. *Note: all imaged individuals have provided photographic consent.*

Neonates were visited at their home address between 24-192 hours post-delivery. Neonates originated from a wide area of villages across the Kombo region, the closest of which was from fifty metres away from the hospital, with the furthest coming from Faraba Banta (41.2 km away). Gambian homes were generally made of concrete with open windows and tin rooftops. Occupancies were shared between large family groups, as is common in the region. At each visit, a physical examination of the neonate was completed, followed by the collection of a venous blood sample, if the neonate was born full-term, normal birthweight. For further details of community visit protocols, see **Chapter 4**.<sup>6</sup>



**Figure 2.5** Community study visits involved a complete review of systems and the collection of neonatal vital signs data. *Note: all imaged individuals have provided photographic consent.*

### **2.1.5 Study Sites: Laboratory Work (Kanifing General Hospital and MRCG Keneba Laboratory)**

During the study site setup process, we became aware that laboratory facilities on-site at the hospital would be required. Kanifing General Hospital provided our study team with a disused storeroom, which we converted into a study-specific laboratory. This fully functioning laboratory contained temperature control, UPS, internet, refrigeration, locked storage and sample processing equipment (i.e. centrifuge, full blood count machine, sample rollers, pipettes). Twelve Kanifing General Hospital and three MRCG laboratory technicians



provided round-the-clock laboratory service. This design ensured the implementation of the study led to direct capacity building in the hospital. Full haematological analyses, serum separation and sample storage were performed in this laboratory. On completing the sample collection phase of the Neolnnate Study, samples were transported to MRCG Keneba field station (Kiang region) on dry ice. Secondary laboratory analysis including glucose-6-phosphate dehydrogenase deficiency testing (G6PD), biochemical analysis (serum ferritin, serum iron, unbound iron-binding capacity (UIBC), soluble transferrin receptor (sTfR), transferrin, C-reactive protein (CRP), haptoglobin, and alpha-1-acid glycoprotein (AGP)) and hepcidin ELISA testing were performed on a batch basis by two study staff members.



**Figure 2.6** Laboratory bench-top work during PhD study. Primary sample processing was completed in a purpose-built renovated laboratory at Kanifing General Hospital for Neolnnate study (*right*). MRC Keneba Microbiology Laboratory (*bottom left*) was the location for secondary sample processing (hepcidin ELISA - *top left*) and analysis.

## 2.2 References

- 1 World Bank Country Profile | Data. Available at:  
<https://data.worldbank.org/country/gambia> (accessed Dec 9, 2019).
- 2 World Bank Country Profile | World Development Indicators. Available at:  
[https://databank.worldbank.org/views/reports/reportwidget.aspx?Report\\_Name=CountryProfile&Id=b450fd57&tbar=y&dd=y&inf=n&zm=n&country=GMB](https://databank.worldbank.org/views/reports/reportwidget.aspx?Report_Name=CountryProfile&Id=b450fd57&tbar=y&dd=y&inf=n&zm=n&country=GMB) (accessed Dec 9, 2019).
- 3 The Gambia Bureau of Statistics (GBOS) and ICF International 2014. The Gambia Demographic and Health Survey 2013. Banjul, The Gambia, and Rockville, Maryland, USA: GBOS and ICF International.
- 4 Gambian Government - Gambian Bureau of Statistics (GBOS) | Statistical Abstract 2016. Available at: <http://www.gbos.gov.gm/uploads/coredocs/Statistical-Abstract-2016.pdf>. (accessed Dec 9, 2019).
- 5 Blencowe H, Cousens S, Jassir FB, *et al*. National, regional, and worldwide estimates of stillbirth rates in 2015, with trends from 2000: A systematic analysis. *Lancet Glob Heal* 2016;**4**:e98–108.
- 6 Cross JH, Jarjou O, Mohammed NI, Prentice AM, Cerami C. Neonatal iron distribution and infection susceptibility in full term, preterm and low birthweight babies in urban Gambia: study protocol for an observational study. *Gates Open Res* 2019;**3**:1469.

## Chapter 3 - Hepcidin, serum iron and transferrin saturation in full term and premature infants during the first month of life: A review of existing evidence in humans (Review)

---

### Summary of Chapter

**BACKGROUND:** Iron sequestration is a fundamental aspect of innate immunity. Evidence suggests that neonates actively regulate their iron distribution at birth and in early postnatal life, in order to sequester iron away from microbial pathogens.

**OBJECTIVE:** To review existing evidence of serum iron, transferrin saturation and hepcidin levels in full-term and premature infants during the first month of life.

**METHODS:** We reviewed literature retrieved from PubMed and Ovid Medline containing data on umbilical cord and venous blood concentrations of hepcidin, serum iron and transferrin saturation in human neonates from 0-1 month of age. After the standardisation of the hepcidin values based on inter-assay cross-calibration studies, weighted mean averages were produced for cord and venous blood using available data. Similar weighted means were produced for serum iron and TSAT.

**RESULTS:** Data from 59 studies were used to create reference ranges for full-term neonates over the first month of life. In full-term neonates, venous blood hepcidin increases 2-3-fold over the first month of life (to reach 61.1 ng/mL; CI: 20.1-102.0 ng/mL) compared to umbilical cord blood (29.7 ng/mL; CI: 21.1-38.3 ng/mL). Cord blood has high levels of serum iron (28.5 µmol/L; CI: 26.0-31.1 µmol/L) and TSAT (51.7%; CI: 46.5-56.9%). Following a short-lived immediate postnatal hypoferremia, iron and TSAT rebounded to approximately

half the levels in the cord by the end of the first month. There was insufficient data to formulate reference ranges for preterm babies.

**CONCLUSION:** Evidence shows that full-term neonates experience an early postnatal hypoferrremia as an innate immune defence, probably mediated by hepcidin. It is unclear whether preterm babies are able to elicit the same defence.

## RESEARCH PAPER COVER SHEET - REVIEW

---

Please note that a cover sheet must be completed for each research paper included within a thesis.

### **SECTION A – Student Details**

<b>Student ID Number</b>	LSH158152	<b>Title</b>	Mr.
<b>First Name(s)</b>	James Henry		
<b>Surname/Family Name</b>	Cross		
<b>Thesis Title</b>	Iron and Infection: Neonatal Iron Transition		
<b>Primary Supervisor</b>	Dr Carla Cerami		

If the Research Paper has previously been published please complete Section B, if not please move to Section C.

### **SECTION B – Paper already published**

Where was the work published?

When was the work published?

If the work was published prior to registration for your research degree, give a brief rationale for its inclusion

Have you retained the copyright for the work?*	Choose an item.	Was the work subject to academic peer review?	Choose an item.
--	-----------------	---	-----------------

\*If yes, please attach evidence of retention. If no, or if the work is being included in its published format, please attach evidence of permission from the copyright holder (publisher or other author) to include this work.

### **SECTION C – Prepared for publication, but not yet published**

Where is the work intended to be published? Current Developments in Nutrition

Please list the paper's authors in the intended authorship order: James H. Cross, Andrew M. Prentice and Carla Cerami.

Stage of publication Undergoing revision

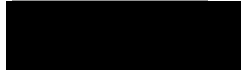
### **SECTION D – Multi-authored work**

For multi-authored work, give full details of your role in the research included in the paper and in the preparation of the paper. (Attach a further sheet if necessary)

James H. Cross is a PhD student with MRCG at LSHTM. He contributed to the literature search, the data analysis and the drafting of the manuscript.

### **SECTION E**

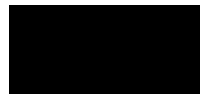
**Student Signature**



**Date**

08/09/2019

**Supervisor Signature**



**Date**

08/09/2019

**Title**

**Hepcidin, serum iron and transferrin saturation in full term and premature infants during the first month of life: A review of existing evidence in humans**

**Authors**

James H Cross<sup>1</sup>, Andrew M Prentice<sup>1</sup>, Carla Cerami<sup>1\*</sup>.

**Affiliations**

<sup>1</sup>Medical Research Council Unit The Gambia at the London School of Hygiene & Tropical Medicine, Atlantic Boulevard, Fajara, P.O. Box 273, Banjul, The Gambia

**\*Corresponding Author**

Dr Carla Cerami, email: [ccerami@mrc.gm](mailto:ccerami@mrc.gm)

Medical Research Council Unit The Gambia at the London School of Hygiene & Tropical Medicine, Atlantic Boulevard, Fajara, P.O. Box 273, Banjul, The Gambia (+220 787 5756)

**Abbreviations:**

AGA	Appropriate for gestational age
CI	Confidence Interval
CRP	C-reactive protein
DFID	Department for International Development
DMT-1	Divalent metal transporter 1
EIA	Enzyme immunoassay
ELISA	Enzyme-linked immunosorbent assays
EPO	Erythropoietin
FTB	Full-term birth

IL-6	Interleukin 6
IQR	Interquartile range
MRC	Medical Research Council
MS	Mass spectrometry
NHNES	National Health and Nutrition Examination Survey
PTB	Preterm birth
SD	Standard Deviation
SGA	Small for gestational age
sTfR	Soluble transferrin receptor
TFR1	Transferrin receptor 1
TLR4	Toll-like receptor 4
TSAT	Transferrin saturation
WHO	World Health Organisation

**Funding:** Bill & Melinda Gates Foundation (OPP1152353) and UK Medical Research Council.

**Competing Interests:** The authors declare that they have no competing interests.



### 3.1 ABSTRACT

**BACKGROUND:** Iron sequestration is a fundamental aspect of innate immunity. Evidence suggests that neonates actively regulate their iron distribution at birth and in early postnatal life, in order to sequester iron away from microbial pathogens.

**OBJECTIVE:** To review existing evidence of serum iron, transferrin saturation and hepcidin levels in full-term and premature infants during the first month of life.

**METHODS:** We reviewed literature retrieved from PubMed and Ovid Medline containing data on umbilical cord and venous blood concentrations of hepcidin, serum iron and transferrin saturation in human neonates from 0-1 month of age. After the standardisation of the hepcidin values based on inter-assay cross-calibration studies, weighted mean averages were produced for cord and venous blood using available data. Similar weighted means were produced for serum iron and TSAT.

**RESULTS:** Data from 59 studies were used to create reference ranges for full-term neonates over the first month of life. In full-term neonates, venous blood hepcidin increases 2-3-fold over the first month of life (to reach 61.1 ng/mL; CI: 20.1-102.0 ng/mL) compared to umbilical cord blood (29.7 ng/mL; CI: 21.1-38.3 ng/mL). Cord blood has high levels of serum iron (28.5  $\mu$ mol/L; CI: 26.0-31.1  $\mu$ mol/L) and TSAT (51.7%; CI: 46.5-56.9%). Following a short-lived immediate postnatal hypoferremia, iron and TSAT rebounded to approximately half the levels in the cord by the end of the first month. There was insufficient data to formulate reference ranges for preterm babies.

**CONCLUSION:** Evidence shows that full-term neonates experience an early postnatal hypoferremia as an innate immune defence, probably mediated by hepcidin. It is unclear whether preterm babies are able to elicit the same defence.

**WORD COUNT ABSTRACT: 260/300**

**KEYWORDS:** Nutritional immunity, host-pathogen interaction, hepcidin, neonates, hypoferremia, transferrin, serum iron

## 3.2 INTRODUCTION

### Iron homeostasis during pregnancy

Three important mediators of hepcidin synthesis: iron status, inflammation, and erythropoiesis, are all altered during pregnancy.<sup>1–4</sup> Iron demand on the mother increases significantly to support expanded maternal erythropoiesis and iron requirements of the growing fetus.<sup>5–9</sup> During pregnancy, the placenta transfers ~270mg of iron from the mother to the fetus via the placenta.<sup>10,11</sup> Syncytiotrophoblasts in the placental villi take up transferrin-bound iron from the maternal circulation by endocytosis via transferrin receptor 1 (TFR1) (**Figure 1**).<sup>12–14</sup> As reviewed in Fisher *et al.*,<sup>15</sup> iron is released from TFR1 and transferred from clathrin-coated vesicles into the syncytiotrophoblast cytoplasm by DMT-1,<sup>16</sup> Zrt/Irt-like protein ZIP8,<sup>17</sup> and ZIP14,<sup>18</sup> collectively. Ferroportin transports iron out of placental syncytiotrophoblasts, and then ceruloplasmin, hephaestin, and zyklopen may all play a role<sup>19–21</sup> in helping the iron pass through the endothelium to reach the fetal circulation and bind to fetal transferrin.<sup>13</sup>

### Maternal control of fetal and early neonatal iron metabolism

Increases in maternal and fetal iron absorption occur in the second and third trimester,<sup>22,23</sup> when maternal hepcidin decreases to trigger increased duodenal iron absorption,<sup>24</sup> splenic macrophage iron recycling, and the release of maternal hepatic iron stores.<sup>25–27</sup> The resulting increased circulating maternal iron is then freely available for transfer to the fetus. Factors that are thought to contribute to maternal hepcidin suppression in the second and third trimester include maternal iron deficiency, erythropoiesis in the mother or fetus,<sup>27</sup> oestrogen,<sup>28</sup> and progesterone receptor membrane component-1.<sup>29</sup> Conflicting evidence now exists as to whether pregnancy-induced plasma dilution may also play a role.<sup>15,30</sup>

### **Fetal control of fetal and early neonatal iron metabolism**

Eighty percent of all the iron transferred from the mother to the fetus occurs in the last trimester.<sup>31</sup> An illustration of the fetal demand for iron (amounting to 1.6-2.0 mg/kg per day<sup>32</sup>) is that umbilical cord blood contains a higher serum iron concentration than in the maternal circulation and at delivery babies have higher total body iron per kilo than that measured in their mothers or in healthy adults.<sup>33-44</sup> This pattern is seen even in anaemic mothers and their babies.<sup>32,43,45,46</sup> The relative roles of maternal and fetal hepcidin levels in controlling placental iron transport are unclear and may change during the course of gestation.<sup>25,26,30,42,45,47-55</sup> As iron becomes more available in the last months of pregnancy, the fetus synthesizes hepcidin probably to control the rate of placental iron transfer and thereby to protect itself from iron-overload.<sup>15,30,56</sup> Evidence showing the importance of fetal hepcidin includes: 1) umbilical cord hepcidin concentrations at birth are higher than maternal levels before and during delivery<sup>26,44,57-60</sup> and 2) in pregnancies with multiple gestations, differences in cord hepcidin between siblings explained a greater fraction of variability in cord hemoglobin, serum ferritin, sTfR, and EPO than maternal hepcidin levels.<sup>49</sup>

### **Placental control of fetal and early neonatal iron metabolism**

The placenta may also independently regulate iron transfer to the fetus in some scenarios.<sup>61</sup> A reduction of ferroportin expression on the apical fetal-facing membrane of placental syncytiotrophoblasts during maternal iron deficiency, in addition to increased expression of TFR1 on the maternal-facing side supports this hypothesis.<sup>30</sup> Sangkhae *et al.* propose that during maternal iron deficiency, iron is held in the placenta to ensure that its metabolic homeostasis is maintained. Placental protein synthesis and critical transfer mechanisms can then continue, ensuring the more detrimental condition of placental dysfunction does not occur. These findings were observed in murine and *in vivo* human trophoblast models, but not in respect to the human pregnancies analysed.<sup>30</sup>

### **Impact of labour and delivery on hepcidin**

Childbirth is an intensely stressful event. Inflammatory pathways (including IL-6) are induced at the onset of human labor, even in the absence of intrauterine infection.<sup>62–69</sup> Initiating stimuli for IL-6 production and release could involve the endocrine events of labor,<sup>68–70</sup> mechanical distension of the membranes and cervix (smooth muscle),<sup>62,70–73</sup> placental hypoxia and/or hypo-perfusion,<sup>70,74</sup> fetal hypoxia-acidemia,<sup>75</sup> pain<sup>76</sup> or exposure to infective agents.<sup>67,69,70,77</sup> The production of IL-6 leads to an increase in hepcidin levels along with a massive influx of immune cells (predominantly neutrophils) into the cervix, decidua, myometrium, chorioamnionic membranes and amniotic fluid.<sup>68,78</sup> This further exacerbates the rise in IL-6 and other cytokines.<sup>76,79</sup> The increase in post-delivery maternal hepcidin concentrations is larger with caesarean section deliveries (5.5-fold increase) as compared to standard vaginal deliveries (3-fold increase).<sup>80</sup> This is most likely due to the surgical procedure and the subsequent inflammation. Similar increases in serum hepcidin are seen postoperatively during other abdominal surgeries.<sup>81</sup> The effect of this maternal rise in hepcidin before, during and immediately after childbirth on the late fetal/early neonatal iron status is unknown, though like IL-6,<sup>82</sup> hepcidin is not thought to cross the placenta.<sup>83</sup>

### **Effects of infection on neonatal serum hepcidin levels**

Intra-amniotic infections can cause an increase in fetal hepcidin.<sup>84</sup> Multiple studies have documented an association between chorioamnionitis, perinatal acidosis and neonatal sepsis with high umbilical cord hepcidin concentrations.<sup>84–89</sup> For example, an extremely high cord concentration (437.6 ng/mL) was found in a neonate with confirmed *Enterococcus faecalis* early-onset sepsis.<sup>87</sup> Similarly, very-low birth weight, premature neonates with late-onset culture-confirmed sepsis, exhibit elevated levels of hepcidin.<sup>90</sup> Nevertheless, despite the well-documented regulatory pathways of infection and inflammation on iron regulation, it is important to note that multiple publications have shown a lack of correlation between hepcidin, IL-6 and CRP in sick neonates.<sup>87,91</sup> This is likely due to differences in the

biochemical kinetics of these molecules. IL-6 concentrations spike very early in the course of perinatal infection, whereas the rise of CRP is delayed.

### **Standardising hepcidin measurements**

Multiple assays, including mass spectrometry (MS) and immunochemistry ELISA methods, are available to quantify hepcidin in various body fluids (urine, serum and plasma).<sup>92</sup>

However, in the studies included in this review, none of these methods are calibrated using the same standards and, as a result, there are significant differences in hepcidin values between studies.<sup>93,94</sup>

In 2016, Van der Vorm *et al.* harmonized many of the available hepcidin ELISA assays using native, lyophilized plasma with cryoprotectant as a commutable candidate reference material.<sup>93</sup> Linear equations were formulated to standardize the hepcidin assays.<sup>93</sup> These equations can now be used to conduct post-hoc standardization of non-calibrated test results, aiding the retrospective comparison of data from previous publications. We have used these equations in this review to generate standardized hepcidin values (**Supplementary Table 1**). The production of standardized reference material, which was refined in 2019, is available for purchase allowing hepcidin measurements to be standardized in all laboratories.<sup>94</sup>

## **3.3 METHODS**

In March 2019, we reviewed the literature searching two databases: PubMed and Ovid Medline with no restrictions on language or the year of publication. The original search was for human studies only. Corresponding authors of extracted publications were not contacted. One individual carried out the inclusion/exclusion process of the retrieved studies, and there was no assessment of bias or the quality of studies as seen in a systematic review process.

**Table 1** displays the search strategy used. **Figure 2** shows the flow diagram of the literature search. The search generated publications containing data on cord and venous concentrations of hepcidin, serum iron and transferrin saturation in the neonatal period. Studies that analyzed healthy neonates were included. Mean, median or range of the gestational age of the study population was a requirement for inclusion. Neonates >37 weeks at delivery were regarded as full-term neonates (FTB). Studies or study groups with a gestational <37 weeks were classed as premature (PTB) neonates. Retrieved publications had to report a mean time of bleed 0-720 hours post-delivery to be analysed. Mean (SD or 95% CI), or median (range, IQR, or 95% CI) data were extracted from the included publications. Studies reporting mean (95% CI) were included in the calculation of weighted means (95% CI) and the associated **Figures 2-5**. Reference ranges for adults and children were presented for comparison.<sup>95,96</sup> Many retrieved publications did not stratify results by birthweight; as a result, this variable was not recorded in **Tables 2-4**. Publications were not stratified by sample type (serum or plasma) due to the overall lack of studies. If multiple publications on the same study population were retrieved, only one was included in the analysis. This was the case with *Ervasti et al.*<sup>97,98</sup> and *Sweet et al.*,<sup>99</sup> as previously mentioned in *Lorenz et al.*<sup>100</sup>

The standardization of hepcidin values using different ELISA assays was performed using the slopes and intercepts from *Van der Vorm et al.*<sup>93</sup> This was performed for studies that used ELISA test kits from DRG (hepcidin-25 (human) EIA Kit, DRG, USA), Bachem (hepcidin-25 EIA Kit, Bachem, USA) and Intrinsic Lifesciences (Intrinsic Hepcidin ELISA Kit, Intrinsic Lifesciences, USA). It was not possible to standardize hepcidin values acquired using the ELISA from Hangzhou Eastbiopharm (Hangzhou Eastbiopharm Co. Ltd. Hangzhou, Zhejiang, China) and mass spectroscopy (MCProt Biotechnology, Kanazawa, Japan), used in *Basu et al.*<sup>52</sup> and *Ichinomiya et al.*,<sup>85</sup> respectively. Prohepcidin was not included in the analysis as it is a poor proxy for biochemically active hepcidin-25.<sup>101–106</sup>

The software Stata IC version 15 (StataCorp LP, College Station, Texas, USA) and R (R: A Language and Environment for Statistical Computing, R Foundation for Statistical Computing, 2020, <https://www.R-project.org>) were used to analyse data. To calculate the confidence interval (CI) around the weighted mean, the weighted variance was calculated using the *wtd.var* function from the R package Hmisc. The standard error derived from this weighted variance was then used to calculate the t-statistic (i.e. weighted mean divided by weighted standard error), from which the 95% CI was derived. GraphPad Prism version 8 (GraphPad Software, San Diego, California, USA) software was used to produce the graphical representation of the results.

### 3.4 RESULTS

The initial search of two electronic databases for three different iron markers yielded 13,931 publications. After the exclusion of duplicated studies and selection criteria filtering, 20 publications were included in the analysis for hepcidin, 23 publications for TSAT and 51 publications for serum iron. Many of these studies were found to contain information on multiple parameters of interest. Overall, we identified 59 publications containing data on hepcidin, serum iron or TSAT in FTB neonates. Sixteen publications were found to contain data on PTB neonates.

In publications detailing the effects of cord clamping interventions, all retrieved cord blood values were from groups that underwent 60 seconds of delayed cord clamping. This is consistent with current WHO policy.<sup>107</sup> Cord blood weighted mean values are generated in **Tables 2-7**, and are represented by a dashed line in **Figures 3-5** and  $\alpha$  (95% CI) in **Figures 6-8**.

## Hepcidin

Standardized weighted mean umbilical cord blood hepcidin levels were higher in FTB neonates (29.7 ng/mL; CI: 21.1-38.3 ng/mL) vs PTB neonates (8.4 ng/mL; CI: 2.0-14.7 ng/mL) (**Figure 3A** and **3B** and **Tables 2** and **3**). Full-term cord blood hepcidin levels were 2-fold higher than in adult male (13.1 ng/mL; CI: 1.4-43.2 ng/mL) and female (10.6 ng/mL; CI: 1.4-43 ng/mL) references ranges (**Table 2**). FTB standardized venous hepcidin levels increased (61.1 ng/mL; CI: 20.1-102.0 ng/mL) over the first four days of life (**Figure 6A**). This trend is unclear for PTB neonates due to the lack of studies. (**Table 3** and **Figure 6B**). No studies were retrieved that assessed post-delivery venous blood samples >77 hours in FTB or >168 hours in PTB.

## TSAT

The weighted mean TSAT in cord blood was higher in FTB neonates (51.7%; CI: 46.5-56.9%) compared to PTB neonates (36.5%; CI: 0.8-72.1%) (**Tables 4** and **5** and **Figure 4**). Cord blood TSAT in FTB neonates was double the reference levels found in adults (23.5%; CI: 12-38.8%) and children aged 1-5 years (19.4%; CI: 8.2-32.9%) (**Table 4**). The weighted mean average of TSAT decreased 2-fold from cord blood to venous blood in FTB neonates (down to 25.2%; CI: 20.1-30.3%) (**Figure 7A**). This hypoferremic response in FTB neonates was followed by a steady increase from 21.8% (CI: 18.8-24.7%) to 44.2% (CI: 32.1-57.8%). No trend was identifiable in PTB neonates due to the lack of data (**Table 5** and **Figure 7B**).

## Serum Iron

Unlike TSAT values, serum iron levels in cord blood were higher in PTB (46.8 µmol/L; CI: 29.7-63.8 µmol/L) neonates compared to FTB neonates (28.5 µmol/L; CI: 26.0-31.1 µmol/L) (**Figure 5**). Like TSAT, a similar 2-fold decrease in the weighted mean average of venous blood compared to cord blood is seen in FTB (13.8 µmol/L; CI: 10.8-16.9 µmol/L) (**Table 6**), and PTB neonates (16.2 µmol/L; CI: 15.3-17.0 µmol/L) (**Table 7**). **Figure 8** suggests that



after the initial reduction (in the first 48 hours of life), levels of serum iron remain consistent over the first month of life in PTB (**B**) and FTB neonates (**A**). Serum iron was lowest between 0-48 hours post-delivery (**Table 6** and **Table 7**).

### 3.5 DISCUSSION

#### Standardization of hepcidin

To our knowledge, this is the first review to retrospectively compare serum hepcidin concentrations between studies, using post-hoc standardized values to produce calculated weighted mean averages in umbilical cord and venous blood. Pre-standardized hepcidin values in cord and peripheral blood share good correlation between methods. However, absolute values vary considerably between the ELISA test kits used (**Table 2-3**). A trend of low values is associated with the use of DRG ELISA kits, and high values linked with the use of Intrinsic Lifesciences and Bachem ELISA kits.

#### Hypoferremia in FTB neonates

The weighted mean average for cord blood hepcidin was calculated using data from 11 studies. Almost all included studies reported a mean value between 11-41ng/mL, apart from Kulik-Rechberger *et al.* This study reported a much higher cord blood hepcidin value (67.9ng/mL; CI: 59.3-76.5ng/mL) as seen in **Figure 3A**. In addition, this study also recorded higher hepcidin levels in venous samples collected at 72 hours (92.9ng/mL; CI: 83.3-102.3ng/mL),<sup>26</sup> compared to those collected by Prentice *et al.* at 77 hours (55.6ng/mL; CI: 47.1-65.5ng/mL).<sup>108</sup>

When all the data are reviewed together (**Figure 6A**), hepcidin increases from within the first 2-11 hours of life<sup>108</sup> and then continues to increase up to 82 hours post-delivery. At all times the hepcidin levels are much higher than those recorded in adults. This excess hepcidin

production may provide a quick, comprehensive and relatively long-lasting (0-3 days) hypoferremic response to aid protection during this vulnerable period<sup>108</sup>. After the first few days, TSAT gradually increases as do serum iron levels, eventually reaching a plateau at approximately 1 month of age.

### **Iron metabolism biomarker data gaps in first month of life in full-term babies**

Gaps in the time course of the concentration of hepcidin, TSAT and serum iron in the first month of life in full-term neonates still exist. This hinders our understanding of neonatal iron metabolism, particularly because hepcidin, TSAT and serum iron are transient and dynamic iron parameters. At the point in which hypoferremia is believed to be maximal, publications detailing the concentration in early (<12 hours) venous samples are lacking in both groups (FTB n=2, PTB n=1). Further research at this time point is required to fully elicit the strength and consistency of this response, as well as understanding the process in greater detail.

### **Lack of data on preterm neonates during the first 24 hours**

After analysis of the current literature, the extent of the role that hypoferremia plays in neonates with a gestational age less than 37 weeks is still unclear. This is primarily due to the limited number of publications documenting hepcidin (n=5), TSAT (n=6) and serum iron (n=13) in the first month of life in preterm neonates. The variability between the studies is vast and further complicated by the complex, intensive and inconsistent care of premature neonates worldwide.

Data analysis of the retrieved publications suggests that preterm neonates have lower cord hepcidin than in full-term neonates, infants and healthy adults. Weighted cord mean values are 3-fold higher in full-term (29.7ng/mL; CI: 21.1-38.3ng/mL) neonates compared to preterm (8.4ng/mL; CI: 2.0-14.7ng/mL) neonates. We speculate that this could be due to very early preterm neonates (<30 weeks' gestation) possessing circulatory monocytes with decreased surface expression of TLR4, lower mRNA expression of TLR4 and reduced cytokine

production.<sup>109</sup> An effect on the production of IL-6 at delivery, might then lead to a reduced ability to stimulate hepcidin expression as suggested in full-term babies.

Our analysis proposes that peripheral venous hepcidin values in preterm neonates increase to 44ng/mL at 168 hours. However, decreases in TSAT between the cord and venous samples are not observed (36.5% to 45.6%). We propose that this is due to a lack of data on TSAT levels in preterm neonates over the first hours of life, potentially due to the complex ethical questions around bleeding preterm neonates so early in postnatal life. This results in the collection of skewed data, focusing only on later time points in the first month of life.

## **Limitations**

The aim of this review was to evaluate our current knowledge on neonatal iron homeostasis in preterm and full-term neonates. As a result of the dearth of publications detailing the parameters of interest during this period, our review has several limitations. Firstly, many studies do not stratify their study groups by gestational age (preterm: <37 weeks, full term: >37 weeks). Subsequently, we have had to assign each study group or population by the mean gestational age. This will result in a reduction of any natural variation potentially caused by gestational age between the reviewed populations. This is also the case with respect to birth weight.

Similarly, the studies on preterm neonates are made up of multiple small sample size subgroups with different gestational ages. Due to the lack of preterm studies, we have had to combine these study groups to formulate weighted means and figures. This in itself, could distort the impact of gestational age on our results, since data from the very early preterm newborns is combined with that from the late preterm neonates.

The retrieval of gestational age was a crucial aspect of the search strategy; however, few studies document the method used. There are large differences in the accuracy of different techniques.<sup>110</sup>

Post-hoc standardization of different hepcidin ELISA kits has, to our knowledge, never been completed before with retrospective data. However, care should be given to the accuracy of the standardized values, as standardization was only possible for DRG, Bachem and Intrinsic Lifesciences ELISA test kits. Studies that used alternative methods<sup>111</sup> were not included in summary statistics.

An essential criterion of inclusion in this publication was that all neonatal data came from healthy newborns. However, documentation of labor practices and postnatal care, along with postnatal medication lack detail in the publications retrieved. Vaginal delivery is commonly referred to as the method of delivery; however, the use of inflammation-inducing forceps or vacuum delivery is not widely reported.

## **Conclusion**

Currently available data suggests that hepcidin, serum iron and TSAT levels for adults and infants are much lower than those found in cord blood and venous blood from neonates during the first month of life. We have strengthened the evidence that full-term neonates possess the ability to produce a hepcidin-mediated hypoferremic response post-delivery. Whether this mechanism is found in PTB neonates is still unclear. This is predominately due to the lack of studies on healthy preterm neonates during the first hours of life. If premature or low birthweight neonates are unable to mount a hypoferremic response, this could enhance their risk of early neonatal infections. Conversely, if the hypoferremic response is seen in both preterm and full-term neonates, it will further support the hypothesis that regulation of iron distribution plays a fundamental role as an innate mechanism of protection against infection.

In summary, serum hepcidin is likely triggered by the inflammatory effect of labor and delivery. We suggest that this intrinsic mechanism of protection protects newborns with immature immune systems to transition from a semi-allogeneic, protected fetal setting to a microbe-rich extrauterine environment.<sup>112,113</sup> Hepcidin-induced hypoferremia then potentially provides a broad action innate bacteriostatic action to invading micro-organisms, when physiological adaption to postnatal life is so critical for survival.

### **Acknowledgements:**

We thank Dr Andrew Armitage for his advice and support in formulating standardised hepcidin values.

### **3.6 REFERENCES**

- 1 Ganz T, Nemeth E. Hepcidin and Disorders of Iron Metabolism. *Annu Rev Med* 2011; **62**: 347–60.
- 2 Choi JW, Pai SH. Change in erythropoiesis with gestational age during pregnancy. *Ann Hematol* 2001; **80**: 26–31.
- 3 Belo L, Santos-Silva A, Rocha S, *et al*. Fluctuations in C-reactive protein concentration and neutrophil activation during normal human pregnancy. *Eur J Obstet Gynecol Reprod Biol* 2005; **123**: 46–51.
- 4 Scholl TO. Iron status during pregnancy: setting the stage for mother and infant. *Am J Clin Nutr* 2005; **81**: 1218S-1222S.
- 5 Barrett JF, Whittaker PG, Williams JG, Lind T. Absorption of non-haem iron from food during normal pregnancy. *BMJ* 1994; **309**: 79–82.
- 6 Milman N. Iron and pregnancy—a delicate balance. *Ann Hematol* 2006; **85**: 559–65.
- 7 Hubbard AC, Bandyopadhyay S, Wojczyk BS, Spitalnik SL, Hod EA, Prestia KA. Effect of dietary iron on fetal growth in pregnant mice. *Comp Med* 2013; **63**: 127–35.

- 8 Lin W-J, Kirksey A. Effects of Different Levels of Dietary Iron on Pregnancy Superimposed upon Growth in the Rat. *J Nutr* 1976; **106**: 543–54.
- 9 Gao G, Liu S-Y, Wang H-J, *et al.* Effects of Pregnancy and Lactation on Iron Metabolism in Rats. *Biomed Res Int* 2015; **2015**: 1–9.
- 10 Bothwell TH. Iron requirements in pregnancy and strategies to meet them. *Am J Clin Nutr* 2000; **72**: 257S-264S.
- 11 Cao C, O'Brien KO. Pregnancy and iron homeostasis: an update. *Nutr Rev* 2013; **71**: 35–51.
- 12 Cao C, Fleming MD. The placenta: the forgotten essential organ of iron transport. *Nutr Rev* 2016; **74**: 421–31.
- 13 Bastin J, Drakesmith H, Rees M, Sargent I, Townsend A. Localisation of proteins of iron metabolism in the human placenta and liver. *Br J Haematol* 2006; **134**: 532–43.
- 14 Georgieff MK, Wobken JK, Welle J, Burdo JR, Connor JR. Identification and Localization of Divalent Metal Transporter-1 (DMT-1) in Term Human Placenta. *Placenta* 2000; **21**: 799–804.
- 15 Fisher AL, Nemeth E. Iron homeostasis during pregnancy. *Am J Clin Nutr* 2017; **106**: 1567S-1574S.
- 16 Georgieff MK, Wobken JK, Welle J, Burdo JR, Connor JR. Identification and Localization of Divalent Metal Transporter-1 (DMT-1) in Term Human Placenta. *Placenta* 2000; **21**: 799–804.
- 17 Gálvez-Peralta M, He L, Jorge-Nebert LF, *et al.* Zip8 zinc transporter: Indispensable role for both multiple-organ organogenesis and hematopoiesis in utero. *PLoS One* 2012; **7**: e36055.
- 18 Hojyo S, Fukada T, Shimoda S, *et al.* The zinc transporter SLC39A14/ZIP14 controls G-protein coupled receptor-mediated signaling required for systemic growth. *PLoS One* 2011; **6**: e18059.
- 19 Chen H, Attieh ZK, Syed BA, *et al.* Identification of zykelopen, a new member of the vertebrate multicopper ferroxidase family, and characterization in rodents and human

- cells. *J Nutr* 2010; **140**: 1728–35.
- 20 Guller S, Buhimschi CS, Ma YY, *et al*. Placental expression of ceruloplasmin in pregnancies complicated by severe preeclampsia. *Lab Invest* 2008; **88**: 1057–67.
  - 21 Li Y-Q, Bai B, Cao X-X, Yan H, Zhuang G-H. Ferroportin 1 and hephaestin expression in BeWo cell line with different iron treatment. *Cell Biochem Funct* 2012; **30**: 249–55.
  - 22 Hallberg L, Rossander-Hultén L. Iron requirements in menstruating women. *Am J Clin Nutr* 1991; **54**: 1047–58.
  - 23 Guo Y, Zhang N, Zhang D, *et al*. Iron homeostasis in pregnancy and spontaneous abortion. *Am J Hematol* 2019; **94**: 184–8.
  - 24 Whittaker PG, Lind T, Williams JG. Iron absorption during normal human pregnancy: a study using stable isotopes. *Br J Nutr* 1991; **65**: 457–63.
  - 25 Rehu M, Punnonen K, Ostland V, *et al*. Maternal serum hepcidin is low at term and independent of cord blood iron status. *Eur J Haematol* 2010; **85**: 345–52.
  - 26 Kulik-Rechberger B, Kościeszka A, Szponar EEE, *et al*. Hepcidin and iron status in pregnant women and full-term newborns in first days of life. *Ginekol Pol* 2016; **87**: 288–92.
  - 27 Koenig MD, Tussing-Humphreys L, Day J, Cadwell B, Nemeth E. Hepcidin and iron homeostasis during pregnancy. *Nutrients*. 2014; **6**: 3062–83.
  - 28 Hou Y, Zhang S, Wang L, *et al*. Estrogen regulates iron homeostasis through governing hepatic hepcidin expression via an estrogen response element. *Gene* 2012; **511**: 398–403.
  - 29 Li X, Rhee DK, Malhotra R, *et al*. Progesterone receptor membrane component-1 regulates hepcidin biosynthesis. *J Clin Invest* 2015; **126**: 389–401.
  - 30 Sangkhae V, Fisher AL, Wong S, *et al*. Effects of maternal iron status on placental and fetal iron homeostasis. *J Clin Invest* 2020; **130**: 625-640
  - 31 Widdowson EM, Spray CM. Chemical development in utero. *Arch Dis Child* 1951; **26**: 205–14.
  - 32 Fletcher J, Suter PE. The transport of iron by the human placenta. *Clin Sci* 1969; **36**:

- 209–20.
- 33 Awadallah SM, Abu-Elteen KH, Elkarmi AZ, *et al.* Maternal and Cord Blood Serum Levels of Zinc, Copper, and Iron in Healthy Pregnant Jordanian Women. *J Trace Elem Exp Med* 2004; **17**: 1–8.
  - 34 Agrawal RM, Tripathi AM, Agarwal KN. Cord blood haemoglobin, iron and ferritin status in maternal anaemia. *Acta Paediatr Scand* 1983; **72**: 545–8.
  - 35 Lee HS, Kim MS, Kim MH, Kim YJ, Kim WY. Iron status and its association with pregnancy outcome in Korean pregnant women. *Eur J Clin Nutr* 2006; **60**: 1130–5.
  - 36 Prema K. Predictive value of serum copper and zinc in normal and abnormal pregnancy. *Indian J Med Res* 1980; **71**: 554–60.
  - 37 Rougereau A, Goré J, N'diaye M, Person O. Ferritin and iron status in Senegalese women. *Am J Clin Nutr* 1982; **36**: 314–8.
  - 38 Singla PN, Chand S, Agarwal KN. Cord serum and placental tissue iron status in maternal hypoferremia. *Am J Clin Nutr* 1979; **32**: 1462–5.
  - 39 Kocylowski R, Lewicka I, Grzesiak M, *et al.* Evaluation of Mineral Concentrations in Maternal Serum Before and After Birth and in Newborn Cord Blood Postpartum-Preliminary Study. *Biol Trace Elem Res* 2018; **182**: 217–23.
  - 40 Milman N, Ibsen KK, Christensen JM. Serum Ferritin and Iron Status in Mothers and Newborn Infants. *Acta Obstet Gynecol Scand* 1987; **66**: 205–11.
  - 41 Lao TTT, Loong EPLP, Chin RKHK, Lam CWWK, Lam YMM. Relationship between newborn and maternal iron status and haematological indices. *Biol Neonate* 1991; **60**: 303–7.
  - 42 Kelly AM, MacDonald DJ, McDougall AN. Observations on maternal and fetal ferritin concentrations at term. *Br J Obstet Gynaecol* 1978; **85**: 338–43.
  - 43 Choi JW, Kim CS, Pai SH, J.W. C, C.S. K. Erythropoietic activity and soluble transferrin receptor level in neonates and maternal blood. *Acta Paediatr* 2000; **89**: 675–9.
  - 44 Lee S, Guillet R, Cooper EM, *et al.* Prevalence of anemia and associations between



- neonatal iron status, hepcidin and maternal iron status among neonates born to pregnant adolescents. *Pediatr Res* 2016; **79**: 42–8.
- 45 Batey R. Iron and Pregnancy. *Br J Haematol* 1978; **38**: 427–8.
  - 46 Glasser S, Wright C, Heyssel R. Transfer of iron across the placenta and fetal membranes in the rat. *Am J Physiol Content* 1968; **215**: 205–10.
  - 47 Sturgeon P. Studies of iron requirements in infante and children. I. Normal values for serum iron, copper and free erythrocyte protoporphyrin. *Pediatrics* 1954; **13**: 107–25.
  - 48 Wallenburg HC, van Eijk HG. Effect of oral iron supplementation during pregnancy on maternal and fetal iron status. *J Perinat Med* 1984; **12**: 7–12.
  - 49 Ru Y, Pressman EK, Guillet R, Katzman PJ, Vermeylen F, O'Brien KO. Umbilical Cord Hepcidin Concentrations Are Positively Associated with the Variance in Iron Status among Multiple Birth Neonates. *J Nutr* 2018; **148**: 1716–22.
  - 50 Briana DD, Boutsikou T, Baka S, *et al.* Perinatal role of hepcidin and iron homeostasis in full-term intrauterine growth-restricted infants. *Eur J Haematol* 2013; **90**: 37–44.
  - 51 Lee S, Guillet R, Cooper EM, *et al.* Prevalence of anemia and associations between neonatal iron status, hepcidin, and maternal iron status among neonates born to pregnant adolescents. *Pediatr Res* 2016; **79**: 42–8.
  - 52 Basu S, Kumar N, Srivastava R, Kumar A. Maternal and Cord Blood Hepcidin Concentrations in Severe Iron Deficiency Anemia. *Pediatr Neonatol* 2016; **57**: 413–9.
  - 53 Merhi ZO, Seifer DB, Weedon J, *et al.* Circulating vitamin D correlates with serum antimüllerian hormone levels in late-reproductive-aged women: Women's Interagency HIV Study. *Fertil Steril* 2012; **98**: 228–34.
  - 54 Dao MC, Sen S, Iyer C, Klebenov D, Meydani SN. Obesity during pregnancy and fetal iron status: Is hepcidin the link? *J Perinatol* 2013; **33**: 177–81.
  - 55 Puolakka J, Janne O, Vihko R. Evaluation by serum ferritin assay of the influence of maternal iron stores on the iron status of newborns and infants. *Acta Obstet Gynecol Scand Suppl* 1980; **95**: 53–6.
  - 56 Nicolas G, Bennoun M, Porteu A, *et al.* Severe iron deficiency anemia in transgenic

- mice expressing liver hepcidin. *Proc Natl Acad Sci* 2002; **99**: 4596–601.
- 57 Rehu M, Punnonen K, Ostland V, *et al.* Maternal serum hepcidin is low at term and independent of cord blood iron status. *Eur J Haematol* 2010; **85**: 345–52.
- 58 Van Santen S, Wiegerinck ET, Swinkels DW, *et al.* Iron Homeostasis in Mother and Child during Placental Malaria Infection. *Am J Trop Med Hyg* 2011; **84**: 148–51.
- 59 Dao MC, Sen S, Iyer C, Klebenov D, Meydani SN. Obesity during pregnancy and fetal iron status: is Hepcidin the link? *J Perinatol* 2013; **33**: 177–81.
- 60 Garcia-Valdes L, Campoy C, Hayes H, Florido J. The impact of maternal obesity on iron status, placental transferrin receptor expression and hepcidin expression in human pregnancy. *Int J Obes* 2015; **39**: 571–8.
- 61 Parrow NL, Fleming RE. The selfishly selfless placenta. *J. Clin. Invest.* 2020; 130: 590–592.
- 62 Santhanam U, Avila C, Romero R, *et al.* Cytokines in normal and abnormal parturition: elevated amniotic fluid interleukin-6 levels in women with premature rupture of membranes associated with intrauterine infection. *Cytokine* 1991; **3**: 155–63.
- 63 Opsjøn S-L, Wathen NC, Tingulstad S, *et al.* Tumor necrosis factor, interleukin-1, and interleukin-6 in normal human pregnancy. *Am J Obstet Gynecol* 1993; **169**: 397–404.
- 64 Liechty KW, Koenig JM, Mitchell MD, Romero R, Christensen RD. Production of interleukin-6 by fetal and maternal cells in vivo during intraamniotic infection and in vitro after stimulation with interleukin. *Pediatr Res* 1991; **29**: 1–4.
- 65 Hillier SL, Witkin SS, Krohn MA, Watts DH, Kiviat NB, Eschenbach DA. The relationship of amniotic fluid cytokines and preterm delivery, amniotic fluid infection, histologic chorioamnionitis, and chorioamnion infection. *Obstet Gynecol* 1993; **81**: 941–8.
- 66 Austgulen R, Lien E, Liabakk NB, Jacobsen G, Arntzen KJ. Increased levels of cytokines and cytokine activity modifiers in normal pregnancy. *Eur J Obstet Gynecol Reprod Biol* 1994; **57**: 149–55.

- 67 Cox SM, Casey ML, MacDonald PC. Accumulation of interleukin-1 $\beta$  and interleukin-6 in amniotic fluid: A sequela of labour at term and preterm. *Hum. Reprod. Update*. 1997; **3**: 517–27.
- 68 Christiaens I, Zaragoza DB, Guilbert L, Robertson SA, Mitchell BF, Olson DM. Inflammatory processes in preterm and term parturition. *J. Reprod. Immunol*. 2008; **79**: 50–7.
- 69 Keelan JA, Blumenstein M, Helliwell RJA, Sato TA, Marvin KW, Mitchell MD. Cytokines, prostaglandins and parturition - A review. *Placenta* 2003; **24**: 33-46
- 70 Bowen JM, Chamley L, Mitchell MD, Keelan JA. Cytokines of the placenta and extra-placental membranes: Biosynthesis, secretion and roles in establishment of pregnancy in women. *Placenta*. 2002; **23**: 239–56.
- 71 El Maradny E, Kanayama N, Halim A, Maehara K, Terao T. Stretching of fetal membranes increases the concentration of interleukin- 8 and collagenase activity. *Am J Obstet Gynecol* 1996; **174**: 843–9.
- 72 Maehara K, Kanayama N, Maradny EE, Uezato T, Fujita M, Terao T. Mechanical stretching induces interleukin-8 gene expression in fetal membranes: a possible role for the initiation of human parturition. *Eur J Obstet Gynecol Reprod Biol* 1996; **70**: 191–6.
- 73 Sivarajasingam SP, Imami N, Johnson MR. Myometrial cytokines and their role in the onset of labour. *J. Endocrinol*. 2016; **231**: R101–19.
- 74 Pierce BT, Pierce LM, Wagner RK, *et al*. Hypoperfusion causes increased production of interleukin 6 and tumor necrosis factor  $\alpha$  in the isolated, dually perfused placental cotyledon. *Am J Obstet Gynecol* 2000; **183**: 863–7.
- 75 Chiesa C, Pellegrini G, Panero A. Umbilical cord interleukin-6 levels are elevated in term neonates with perinatal asphyxia. *Eur J Clin Invest* 2003; **33**: 352–8.
- 76 Saito S, Kasahara T, Kato Y, Ishihara Y, Ichijo M. Elevation of amniotic fluid interleukin 6 (IL-6), IL-8 and granulocyte colony stimulating factor (G-CSF) in term and preterm parturition. *Cytokine* 1993; **5**: 81–8.

- 77 Gotsch F, Romero R, Kusanovic JP, *et al.* The fetal inflammatory response syndrome. *Clin. Obstet. Gynecol.* 2007; **50**: 652–83.
- 78 Osman I, Young A, Ledingham MA, *et al.* Leukocyte density and pro-inflammatory cytokine expression in human fetal membranes, decidua, cervix and myometrium before and during labour at term. *Mol. Hum. Reprod.* 2003; **9**: 41–5.
- 79 Stallmach T, Hebisch G, Joller-Jemelka HI, Orban P, Schwaller J, Engelmann M. Cytokine production and visualized effects in the feto-maternal unit. Quantitative and topographic data on cytokines during intrauterine disease. *Lab Invest* 1995; **73**: 384–92.
- 80 Gyarmati B, Szabó E, Szalay B, *et al.* Serum maternal hepcidin levels 3 days after delivery are higher compared to those measured at parturition. *J Obstet Gynaecol Res* 2011; **37**: 1620–4.
- 81 Park KH, Sawada T, Kosuge T, *et al.* Surgical Inflammation Induces Hepcidin Production after Abdominal Surgery. *World J Surg* 2012; **36**: 800–6.
- 82 Aaltonen R, Heikkinen T, Hakala K, Laine K, Alanen A. Transfer of Proinflammatory Cytokines Across Term Placenta. *Obstet Gynecol* 2005; **106**: 802–7.
- 83 Hoppe M, Hulthén L, Samuelson G. Is cord blood hepcidin influenced by the low-grade acute-phase response occurring during delivery? A small-scale longitudinal study. *J. Matern. Neonatal Med.* 2018; **32**: 2166-2172
- 84 Tabbah SM, Buhimschi CS, Rodewald-Millen K. Hepcidin, an Iron Regulatory Hormone of Innate Immunity, is Differentially Expressed in Premature Fetuses with Early-Onset Neonatal Sepsis. *Am J Perinatol* 2018; **35**: 865-872
- 85 Ichinomiya K, Maruyama K, Inoue T, *et al.* Perinatal Factors Affecting Serum Hepcidin Levels in Low-Birth-Weight Infants. *Neonatology* 2017; **112**: 180–6.
- 86 Wu T-WW, Tabangin M, Kusano R, Ma Y. The utility of serum hepcidin as a biomarker for late-onset neonatal sepsis. *J Pediatr* 2013; **162**: 67–71.
- 87 Lorenz L, Herbst J, Engel C, *et al.* Gestational Age-Specific Reference Ranges of Hepcidin in Cord Blood. *Neonatology* 2014; **106**: 133–9.

- 88 Müller KF, Lorenz L, Poets CF, Westerman M, Franz AR. Hepcidin Concentrations in Serum and Urine Correlate with Iron Homeostasis in Preterm Infants. *J Pediatr* 2012; **160**: 949-953.e2.
- 89 Cizmeci MN, Kara S, Kanburoglu MK, *et al.* Detection of cord blood hepcidin levels as a biomarker for early-onset neonatal sepsis. *Med Hypotheses* 2014; **82**: 310–2.
- 90 Wu T-W, Tabangin M, Kusano R, Ma Y, Ridsdale R, Akinbi H. The Utility of Serum Hepcidin as a Biomarker for Late-Onset Neonatal Sepsis. *J Pediatr* 2013; **162**: 67–71.
- 91 R Franz, G Steinbach, M Kron, F Poh A, Steinbach G, Kron M, Pohlandt F. Interleukin-8: a valuable tool to restrict antibiotic therapy in newborn infants. *Acta Paediatr* 2001; **90**: 1025–32.
- 92 Kroot JJC, Laarakkers CMM, Geurts-Moespot AJ, *et al.* Immunochemical and Mass-Spectrometry-Based Serum Hepcidin Assays for Iron Metabolism Disorders. *Clin Chem* 2010; **56**: 1570–9.
- 93 van der Vorm LN, Hendriks JCM, Laarakkers CM, *et al.* Toward Worldwide Hepcidin Assay Harmonization: Identification of a Commutable Secondary Reference Material. *Clin Chem* 2016; **62**: 993–1001.
- 94 Diepeveen LE, Laarakkers CMM, Martos G, *et al.* Provisional standardization of hepcidin assays: creating a traceability chain with a primary reference material, candidate reference method and a commutable secondary reference material. *Clin Chem Lab Med* 2019; **57**: 864–72.
- 95 U.S. Centers for Disease Control. National Report on Biochemical Indicators of Diet and Nutrition in the U.S. Population 1999-2002 - National Health and Nutrition Examination Survey. [https://www.cdc.gov/nutritionreport/99-02/pdf/nr\\_ch3.pdf](https://www.cdc.gov/nutritionreport/99-02/pdf/nr_ch3.pdf) (accessed Oct 26, 2019).
- 96 Radboudumc. Reference values. <http://www.hepcidinanalysis.com/provided-service/reference-values/> (accessed Oct 26, 2019).
- 97 Ervasti M, Sankilampi U, Heinonen S, *et al.* Novel red cell indices indicating reduced availability of iron are associated with high erythropoietin concentration and low ph

- level in the venous cord blood of newborns. *Pediatr Res* 2008; **64**: 135–40.
- 98 Ervasti M, Sankilampi U, Heinonen S, Punnonen K. Early signs of maternal iron deficiency do not influence the iron status of the newborn, but are associated with higher infant birthweight. *Acta Obstet Gynecol Scand* 2009; **88**: 83–90.
  - 99 Sweet DG, Savage G, Tubman TR, *et al.* Study of maternal influences on fetal iron status at term using cord blood transferrin receptors. *Arch Dis Child Fetal Neonatal Ed* 2001; **84**: F40-3.
  - 100 Lorenz L, Peter A, Poets CF, Franz AR. A review of cord blood concentrations of iron status parameters to define reference ranges for preterm infants. *Neonatology*. 2013; **104**: 194–202.
  - 101 Balogh Á, Szabó M, Kelen D, *et al.* Prohepcidin levels during human perinatal adaptation. *Pediatr Hematol Oncol* 2007; **24**: 361-8
  - 102 Frazer DM, Anderson GJ. Hepcidin compared with prohepcidin: An absorbing story. *Am J Clin Nutr* 2009; **89**: 475–6.
  - 103 Slomka A, Korbai P, Piekus N, *et al.* The use of cluster and principal component analysis in the estimation of iron status in term newborns. *J Matern Neonatal Med* 2013; **26**: 482–6.
  - 104 Sasu BJ, Li H, Rose MJ, Arvedson TL, Doellgast G, Molineux G. Serum hepcidin but not prohepcidin may be an effective marker for anemia of inflammation (AI). *Blood Cells Mol Dis* 2010; **45**: 238–45.
  - 105 Beirão I, Almeida S, Swinkels D, *et al.* Low serum levels of prohepcidin, but not hepcidin-25, are related to anemia in familial amyloidosis TTR V30M. *Blood Cells Mol Dis*; **41**: 175–8.
  - 106 Young MF, Glahn RP, Ariza-Nieto M, *et al.* Serum hepcidin is significantly associated with iron absorption from food and supplemental sources in healthy young women. *Am J Clin Nutr* 2009; **89**: 533–8.
  - 107 WHO | Guideline: Delayed umbilical cord clamping for improved maternal and infant health and nutrition outcomes. Geneva: World Health Organisation; 2014.

<https://www.ncbi.nlm.nih.gov/books/NBK310511/>

- 108 Prentice S, Jallow AT, Sinjanka E, *et al.* Hepcidin mediates hypoferremia and reduces the growth potential of bacteria in the immediate post-natal period in human neonates. *Sci Rep* 2019; **9**: e16596
- 109 Forster-Waldl E, Sadeghi K, Tamandl D, *et al.* Monocyte toll-like receptor 4 expression and LPS-induced cytokine production increase during gestational aging. *Pediatr Res* 2005; **58**: 121–4.
- 110 Macaulay S, Buchmann EJ, Dunger DB, Norris SA. Reliability and validity of last menstrual period for gestational age estimation in a low-to-middle-income setting. *J Obstet Gynaecol Res* 2019; **45**: 217–25.
- 111 Basu S, Kumar N, Srivastava R, Kumar A. Effect of Severe Maternal Iron Deficiency Anemia on Neonatal Platelet Indices. *Indian J Pediatr* 2015; **82**: 1091–6.
- 112 de Goffau MC, Lager S, Sovio U, *et al.* Human placenta has no microbiome but can contain potential pathogens. *Nature* 2019; **572**: 329–34.
- 113 Rutayisire E, Huang K, Liu Y, Tao F. The mode of delivery affects the diversity and colonization pattern of the gut microbiota during the first year of infants' life: a systematic review. *BMC Gastroenterol* 2016; **16**: 86.
- 114 NHANES. National Health and Nutrition Examination Survey, 1999-2002 | Iron-Status Indictors. 2003. [https://www.cdc.gov/nutritionreport/99-02/pdf/nr\\_ch3.pdf](https://www.cdc.gov/nutritionreport/99-02/pdf/nr_ch3.pdf)
- 115 Ru Y, Pressman EK, Guillet R, *et al.* Predictors of anemia and iron status at birth in neonates born to women carrying multiple fetuses. *Pediatr Res* 2018; **84**: 199–204.
- 116 Ru Y, Pressman EK, Guillet R, Katzman PJ, Vermeylen F, O'Brien KO. Umbilical Cord Hepcidin Concentrations Are Positively Associated with the Variance in Iron Status among Multiple Birth Neonates. *J Nutr* 2018; **148**: 1716–22.
- 117 Koenig MD, Tussing-Humphreys L, Day J, Cadwell B, Nemeth E. Hepcidin and iron homeostasis during pregnancy. *Nutrients* 2014; **6**: 3062–83.
- 118 Lipiński P, Styś A, Starzyński RR. Molecular insights into the regulation of iron metabolism during the prenatal and early postnatal periods. *Cell. Mol. Life Sci.* 2013;

- 70:** 23–38.
- 119 Sangkhae V, Nemeth E. Placental iron transport: The mechanism and regulatory circuits. *Free Radic. Biol. Med.* 2019; **133**: 254-261.
- 120 Hågå P. Plasma ferritin concentrations in preterm infants in cord blood and during the early anaemia of prematurity. *Acta Paediatr Scand* 1980; **69**: 637–41.
- 121 Sweet DG, Savage GA, Tubman R, Lappin TRJ, Halliday HL. Cord blood transferrin receptors to assess fetal iron status. *Arch Dis Child Fetal Neonatal Ed* 2001; **85**: F46-8.
- 122 Uijterschout L, Domellöf M, Berglund SK, *et al.* Serum hepcidin in infants born after 32 to 37 wk of gestational age. *Pediatr Res* 2016; **79**: 608–13.
- 123 Al-Tawil MM, Abdel-Aal MR, Kaddah MA. A randomized controlled trial on delayed cord clamping and iron status at 3-5 months in term neonates held at the level of maternal pelvis. *J Neonatal Perinatal Med* 2012; **5**: 319-326.
- 124 Andersson O, Hellstrom-Westas L, Andersson D, Domellof M, Hellström-Westas L, Andersson D. Effect of delayed versus early umbilical cord clamping on neonatal outcomes and iron status at 4 months: A randomised controlled trial. *BMJ* 2011; **343**: d7157–d7157.
- 125 Kitajima J, Ohga S, Kinjo T, *et al.* Serum prohepcidin concentrations at birth and 1 month after birth in premature infants. *Pediatr Blood Cancer* 2011; **56**: 267–72.
- 126 Yamada RRT, Leone CCR. Hematological and iron content evolution in exclusively breastfed late-preterm newborns. *Clinics* 2014; **69**: 792–8.
- 127 Lackmann GM, Schnieder C, Bohner J. Gestational Age-Dependent Reference Values for Iron and Selected Proteins of Iron Metabolism in Serum of Premature Human Neonates. *Neonatology* 1998; **74**: 208-13.
- 128 Çelik HT, Yurdakök M, Korkmaz A, Yülit Ş, Yiğit Ş, Yiğit Ş. Serum prohepcidin levels in premature newborns with oxygen radical diseases. *J Matern Neonatal Med* 2015; **28**: 2228–33.
- 129 Patidar S, Shrivastava J, Dwivedi R. Assessment of iron status and red cell



- parameters in healthy full term small for gestational age neonates at birth. *J Clin Neonatol* 2013; **2**: 121.
- 130 Szabo M, Vasarhelyi B, Balla G, Szabo T, MacHay T, Tulassay T. Acute postnatal increase of extracellular antioxidant defence of neonates: The role of iron metabolism. *Acta Paediatr Int J Paediatr* 2001; **90**: 1167–70.
  - 131 Tsuzuki S, Morimoto N, Hosokawa S, Matsushita T. Associations of Maternal and Neonatal Serum Trace Element Concentrations with Neonatal Birth Weight. *PLoS One* 2013; **8**: e75627.
  - 132 Tiker F, Celik B, Tarcan A, *et al.* Serum pro-hepcidin levels and relationships with iron parameters in healthy preterm and term newborns. *Pediatr Hematol Oncol* 2006; **23**: 293–7.
  - 133 Yapakçi E, Tarcan A, Celik B, Ozbek N, Gürakan B. Serum pro-hepcidin levels in term and preterm newborns with sepsis. *Pediatr Int* 2009; **51**: 289–92.
  - 134 Ozkiraz S, Kilicdag H, Gokmen Z, Ecevit A, Tarcan A, Ozbek N. Serum prohepcidin levels and iron parameters in term small-for-gestational-age newborns. *J Matern Fetal Neonatal Med* 2011; **24**: 1437–9.
  - 135 Schiza V, Giapros V, Pantou K, Theocharis P, Challa A, Andronikou S. Serum transferrin receptor, ferritin, and reticulocyte maturity indices during the first year of life in ‘large’ preterm infants. *Eur J Haematol* 2007; **79**: 439–46.

### **Funding:**

The research was undertaken with a research grant provided by the Bill & Melinda Gates Foundation (OPP1152353). The funding agency had no role in the design of this review, and did not have any in the collection, management, analyses or interpretation of the data nor in the preparation, review, or approval of the manuscript. The Nutrition Theme of the MRC Unit The Gambia at LSHTM are supported by core funding MC-A760-5QX00 to the MRC Unit The Gambia/MRC International Nutrition Group by the UK MRC and the UK Department for the International Development (DFID) under the MRC/DFID Concordat agreement.

**Authors' Contributions:**

J.H.C., A.M.P. and C.C. designed the research; J.H.C. conducted the search strategy and analysed data; J.H.C., A.M.P. and C.C. wrote the paper. All authors reviewed the final manuscript prior to submission.

**Consent for Publication:**

Not applicable.

## TABLE LEGENDS:

**Table 1: Literature Search Strategy.** Searches conducted via PubMed and Ovid Medline.

**Table 2: Hepcidin (ng/mL) in full-term newborns over the neonatal period.** Values from Basu *et al*<sup>111</sup> were not standardized because the study used the Hangzhou Eastbiopharm ELISA, which was not part of the Van der Vorm *et al.* analysis.<sup>93</sup> Extracted standard deviations were converted to 95% confidence intervals. Median (IQR or 95% CI) were not included in weighted means. Reference ranges were obtained from Hepcidin Analysis (Radboundumc).<sup>96</sup>

**Table 3: Hepcidin (ng/mL) in preterm newborns over the neonatal period.** Ichinomiya *et al*<sup>65</sup> was not standardized because the study used a mass spectrometry based method that was not part of the Van der Vorm *et al.* analysis.<sup>93</sup> Extracted standard deviations were converted to 95% confidence intervals. Median (IQR or 95% CI) were not included in weighted means. Reference ranges were obtained from Hepcidin Analysis (Radboundumc).<sup>96</sup>

**Table 4: Transferrin saturation (%) in full-term newborns over the neonatal period.** Extracted standard deviations were converted to 95% confidence intervals. Median (IQR or 95% CI) were not included in weighted means. Reference ranges are taken from the National Health and Nutrition Examination Survey, 1999–2000.<sup>114</sup>

**Table 5: Transferrin saturation (%) in preterm newborns over the neonatal period.** Extracted standard deviations were converted to 95% confidence intervals. Median (IQR or 95% CI) were not included in weighted means. AGA group<sup>a</sup> and SGA group<sup>b</sup> can be identified in **Figure 3B** using the superscripted letters. Reference ranges were obtained from the National Health and Nutrition Examination Survey, 1999–2000.<sup>114</sup>

**Table 6: Serum iron (μmol/L) in full-term newborns over the neonatal period.** Extracted standard deviations were converted to 95% confidence intervals. Median (IQR or 95% CI) were not included in weighted means. Reference ranges are generated from the National Health and Nutrition Examination Survey, 1999–2000.<sup>114</sup>

**Table 7: Serum iron (μmol/L) in preterm newborns over the neonatal period.** Extracted standard deviations were converted to 95% confidence intervals. Median (IQR or 95% CI) were not included in weighted means. AGA group<sup>c</sup>, SGA group<sup>d</sup>, 30-36 wks<sup>g</sup> and 24-29 wks<sup>h</sup> can be identified in **Figure 4B** using the superscripted letters. Ru *et al*, 2018<sup>115</sup> is referenced as <sup>e</sup>. Ru *et al*, 2018<sup>116</sup> is referenced as <sup>f</sup>. Reference ranges are generated from the National Health and Nutrition Examination Survey, 1999–2000.<sup>114</sup>

## TABLES:

**Table 1: Literature Search Strategy**

Parameter	Database	Search Strategy
<b>Hepcidin</b>	Ovid Medline	(Human) AND (neonate OR neonates OR infant OR infants OR baby OR babies OR cord OR "umbilical cord".mp.) AND (hepcidin OR prohepcidin.mp.)
	PubMed	(Human) AND (neonate OR neonates OR infant OR infants OR baby OR babies OR cord OR "umbilical cord") AND (hepcidin OR prohepcidin)
<b>TSAT</b>	Ovid Medline	(Human) AND (neonate OR neonates OR infant OR infants OR baby OR babies OR cord OR "umbilical cord".mp.) AND ("transferrin saturation" OR TSAT.mp.)
	PubMed	(Human) AND (neonate OR neonates OR infant OR infants OR baby OR babies OR cord OR "umbilical cord") AND ("transferrin saturation" OR TSAT)
<b>Serum Iron</b>	Ovid Medline	(Human) AND (neonate OR neonates OR infant OR infants OR baby OR babies OR cord OR "umbilical cord".mp.) AND ("serum iron" OR iron.mp.)
	PubMed	(Human) AND (neonate OR neonates OR infant OR infants OR baby OR babies OR cord OR "umbilical cord") AND ("serum iron" OR iron)

**Table 2: Hepcidin concentration (ng/mL) in full-term newborns over the neonatal period.**

Reference	Year	Location	n	Test Type	Type of Sample	Hepcidin (ng/mL)		Standardized Hepcidin (ng/mL)	
						Mean (95% CI)	Median (IQR or 95% CI)	Mean (95% CI)	Median (IQR or 95% CI)
Armitage <i>et al.</i> (VPM)	2019	The Gambia	114	ELISA (Bachem)	Cord (Plasma)		46.0 (31.3-55.1)		29.4 (20-35.2)
Armitage <i>et al.</i> (VA)	2019	The Gambia	193	ELISA (Bachem)	Cord (Plasma)		41.9 (26.3-56.6)		26.8 (16.8-36.1)
Basu <i>et al.</i>	2015	India	15	ELISA (Hangzhou Eastbiopharm)	Cord (Serum)	124.0 (115-133)		N/A	
Briana <i>et al.</i>	2013	Greece	104	ELISA (DRG)	Cord (Serum)		17.85 (4.75-69.2)		24.1 (3.1-44.2)
Cao <i>et al.</i>	2014	USA	57	ELISA (Intrinsic)	Cord (Serum)	131.8 (109-155)		41.7 (34.5-48.9)	
Cao <i>et al.</i>	2016	USA	98	ELISA (Intrinsic)	Cord (Serum)	121.5 (105-138)		38.3 (33.2-43.6)	
Delaney <i>et al.</i>	2019	USA	108	ELISA (Intrinsic)	Cord (Serum)	92.13 (91.9-92.3)		29.2 (29.1-29.2)	
Dosch <i>et al.</i>	2016	USA	47	ELISA (DRG)	Cord (Plasma)	13.4 (11.7-15.1)		17.8 (15.4-20.2)	
Garcia-Valdes <i>et al.</i>	2015	Spain	52	ELISA (DRG)	Cord (Serum)	18.01 (15.1-20.9)		24.3 (20.2-28.4)	
Hoppe <i>et al.</i>	2018	Sweden	15	ELISA (Bachem)	Cord (Serum)		30.5 (21.7-38.8)		19.5 (13.9-24.8)
Kulik-Rechberger <i>et al.</i>	2016	Poland	44	ELISA (DRG)	Cord (Serum)	48.98 (42.9-55.1)		67.9 (59.3-76.5)	
			44		Venous - 72 hours (Serum)	66.79 (60-73.5)		92.9 (83.3-102.3)	
Lee <i>et al.</i>	2016	USA	104	ELISA (Intrinsic)	Cord (Serum)	87.4 (74.4-103)		27.7 (23.6-32.6)	
Lorenz <i>et al.</i>	2014	Germany	100	ELISA (Intrinsic)	Cord (Plasma)		103.9 (61.4-149.2)		32.9 (19.6-47.1)
Prentice <i>et al.</i>	2019	The Gambia	81	ELISA (Bachem)	Cord (Serum)	43.8 (36.8-52.3)		27.9 (23.5-33.4)	
			53		Venous - 6 hours (2-11) (Serum)	79.4 (68.1-92.4)		50.7 (43.5-58.9)	
			21		Venous - 29 hours (26-34) (Serum)	45.9 (36.5-57.8)		29.3 (23.3-36.9)	
			33		Venous - 77 hours (74-82) (Serum)	87.1 (73.8-102.7)		55.6 (47.1-65.5)	
Rehu <i>et al.</i>	2010	Finland	116	ELISA (Intrinsic)	Cord (Serum)	71.6 (60.8-84.4)		22.8 (19.4-26.77)	
Ru <i>et al.</i>	2018	USA	50	ELISA (Bachem)	Cord (Serum)	17 (12-24.2)		10.9 (7.7-15.5)	
Slomka <i>et al.</i>	2013	Poland	54	ELISA (DRG)	Cord (Serum)		18.50 (2.75-35.13)		25 (2.8-48.4)
Young <i>et al.</i>	2011	USA	19	ELISA (Intrinsic)	Cord (Serum)	61.0 (26.4-95.6)		19.4 (8.6-30.3)	
<b>Weighted Mean (Cord)</b>						<b>73.2 (48.1-98.3)</b>	<b>N/A</b>	<b>29.7 (21.1-38.3)</b>	<b>N/A</b>

Weighted Mean (Venous)	72.7 (48.3-97.2)	N/A	61.1 (20.1-102.0)	N/A
Adults (Median (95% CI))	Men: 13.1 (1.4-43.2) ng/mL   Women: 10.6 (1.4-43) ng/mL (hepcidinanalysis.com, 2019)			
Infants (Median (95% CI))	11.9 (3.3-37.7) ng/mL (hepcidinanalysis.com, 2019)			

**Table 3: Hepcidin concentration (ng/mL) in preterm newborns over the neonatal period.**

Reference	Year	Location	n	Test Type	Type of Sample	Study Group	Hepcidin (ng/mL)		Standardized Hepcidin (ng/mL)	
							Mean (95% CI)	Median (IQR or 95% CI)	Mean (95% CI)	Median (IQR or 95% CI)
Delaney <i>et al.</i>	2019	USA	126	ELISA (Bachem)	Cord (Serum)		13.78 (13.6-14)		8.8 (8.7-9)	
Ichinomiya <i>et al.</i>	2017	Japan	92	Mass Spec (MCProt)	Cord (Serum)			7.3 (2.85-16.38)		N/A
Lorenz <i>et al.</i>	2014	Germany	40	ELISA (Intrinsic)	Cord (Plasma)	24-29 wks		26.9 (13.5-63.1)		8.7 (4.5-20.1)
			81	ELISA (Intrinsic)	Cord (Plasma)	30-36 wks		45.9 (24.7-74.5)		14.7 (8.0-23.7)
Ru <i>et al.</i>	2018	USA	92	ELISA (Bachem)	Cord (Serum)		12.1 (9.2-15.7)		7.8 (5.9-10.1)	
Uijterschout <i>et al.</i>	2015	Netherlands	85	ELISA (Bachem)	Venous - 168 hours (Serum)		69.6 (14.6-180.1)		44.4 (9.4-114.8)	
<b>Weighted Mean (Cord)</b>							<b>13.1 (2.4-23.7)</b>	<b>N/A</b>	<b>8.4 (2.0-14.7)</b>	<b>N/A</b>
<b>Weighted Mean (Venous)</b>							<b>69.6 (14.6-180.1)</b>	<b>N/A</b>	<b>44.4 (9.4-114.8)</b>	<b>N/A</b>
<b>Adults (Median (95% CI))</b>							<b>Men: 13.1 (1.4-43.2) ng/mL   Women: 10.6 (1.4-43) ng/mL (hepcidinanalysis.com, 2019)</b>			
<b>Infants (Median (95% CI))</b>							<b>11.9 (3.3-37.7) ng/mL (hepcidinanalysis.com, 2019)</b>			

**Table 4: Transferrin saturation (%) in full-term newborns over the neonatal period.**

Reference	Year	Location	n	Type of Sample	TSAT (%)	
					Mean (95% CI)	Median (IQR or 95% CI or Range)
Al-Tawil <i>et al.</i>	2012	Egypt	90	Venous - 24 hours	25 (24.6-25.4)	
Ali <i>et al.</i>	2016	USA	64	Cord	59.2 (53.9-64.5)	
Anderson <i>et al.</i>	2011	Sweden	162	Venous - 48 hours	23 (21.9-24.1)	
Balogh <i>et al.</i>	2007	Hungary	20	Cord		60.5 (14-90)
			20	Venous - 39 hours (18-114)		22.5 (11-42)
Basu <i>et al.</i>	2015	India	15	Cord	61.8 (54.7-68.9)	
El-Farrash <i>et al.</i>	2012	Egypt	30	Cord	49.5 (42.5-56.5)	
Ervasti <i>et al.</i>	2007	Finland	199	Cord	55 (52.4-57.6)	
Haga <i>et al.</i>	1980	Norway	21	Cord	55 (33.8-76.2)	
Kalem <i>et al.</i>	2019	Turkey	380	Cord	55.87 (54.8-56.9)	
Kelly <i>et al.</i>	1978	Scotland	115	Cord	58.8 (55.6-62)	
Kitajima <i>et al.</i>	2010	Japan	8	Cord		15.1 (8.3-27.5)
			8	Venous - 720 hours		44.2 (32.1-57.8)
Kleven <i>et al.</i>	2007	USA	26	Cord	42 (32.4-51.6)	
Mashako <i>et al.</i>	1991	DRC	166	Cord	32.3 (30.1-34.5)	
Milman <i>et al.</i>	1987	Denmark	74	Cord		48 (32-71)
			47	Venous - 120 hours		33 (21-48)
Prentice <i>et al.</i>	2019	The Gambia	81	Cord	47.6 (43.7-51.5)	
			53	Venous - 6 hours (2-11)	24.4 (21.2-27.6)	
			21	Venous - 29 hours (26-34)	21.8 (18.8-24.7)	
			33	Venous - 77 hours (74-82)	30.9 (26.9-34.8)	
Puolakka <i>et al.</i>	1980	Finland	47	Cord	53 (49-57)	
Rehu <i>et al.</i>	2010	Finland	116	Cord	50.6 (44.5-57.5)	
Rois <i>et al.</i>	1975	USA	26	Cord	61.2 (55.9-66.5)	



Slomka <i>et al.</i>	2013	Poland	49	Cord		58.1 (51.7-73.6)
Yamada <i>et al.</i>	2014	Brazil	21	Cord	47.7 (40.2-55.2)	
			21	Venous - 720 hours	39.8 (34.7-44.9)	
<b>Weighted Mean (Cord)</b>					<b>51.7 (46.5-56.9)</b>	<b>N/A</b>
<b>Weighted Mean (Venous)</b>					<b>25.2 (20.1-30.3)</b>	<b>N/A</b>
<b>Adults (Median (95% CI))</b>					<b>23.5 (12-38.8) % (10-90th percentiles) - NHNES</b>	
<b>Infants (Median (95% CI))</b>					<b>19.4 (8.2-32.9) % (10-90th percentiles) - NHNES</b>	

**Table 5: Transferrin saturation (%) in preterm newborns over the neonatal period.**

Reference	Year	Location	n	Type of Sample	Study Group	TSAT (%)		
						Mean (95% CI)	Median (IQR or 95% CI or Range)	
Celik <i>et al.</i>	2015	Turkey	42	Venous - 648 hours (288-1872)		46.5 (41.2-51.8)		
Haga <i>et al.</i>	1980	Norway	23	Cord	AGA Group <sup>a</sup>	48 (39.8-56.2)		
			6	Cord	SGA Group <sup>b</sup>	41 (23.4-58.6)		
Ichinomiya <i>et al.</i>	2017	Japan	92	Cord		87.2 (68.3-100)		
Kitajima <i>et al.</i>	2010	Japan	13	Cord		64.3 (15.8-88.9)		
			13	Venous - 720 hours		33.2 (17.1-79.5)		
Lackmann <i>et al.</i>	1998	Germany	15	Venous (<1 hour)	<32 wks	39 (5-83)		
			22	Venous (<1 hour)	33-34 wks	36 (7-87)		
			26	Venous (<1 hour)	35-36 wks	31 (13-60)		
Yamada <i>et al.</i>	2014	Brazil	25	Cord		24.8 (18.5-31.1)		
			25	Venous - 720 hours		44.1 (37.3-50.9)		
Weighted Mean (Cord)						36.5 (0.8-72.1)		N/A
Weighted Mean (Venous)						45.6 (30.4-60.9)		N/A
Adults (Median (95% CI))						23.5 (12-38.8) % (10-90th percentiles) - NHNES		
Infants (Median (95% CI))						19.4 (8.2-32.9) % (10-90th percentiles) - NHNES		

**Table 6: Serum iron concentration (µmol/L) in full-term newborns over the neonatal period.**

Reference	Year	Location	n	Type of Sample	Serum Iron (µmol/L)	
					Mean (95% CI or Range)	Median (IQR or 95% CI or Range)
Ahlsten <i>et al.</i>	1989	Sweden	20	Cord	38 (34.9-41.1)	
Ali <i>et al.</i>	2016	USA	64	Cord	26.8 (24.4-29.2)	
Amarnath <i>et al.</i>	1989	USA	15	Cord	24.1 (21-27.2)	
Anderson <i>et al.</i>	2011	Sweden	162	Venous - 48 hours	9.9 (9.5-10.3)	
Armitage <i>et al.</i> (VA)	2019	The Gambia	193	Cord		18.8 (15.4-22.3)
Armitage <i>et al.</i> (VPM)	2019	The Gambia	114	Cord		16.0 (12.7-18.7)
Awadallah <i>et al.</i>	2004	Jordan	92	Cord	20.7 (20.1-21.3)	
Balogh <i>et al.</i>	2007	Hungary	20	Cord		25.5 (8-43)
			20	Venous - 39 hours (18-114)		9.5 (5-20)
Bastida <i>et al.</i>	2000	Spain	70	Cord	41.5 (38.3-44.7)	
Basu <i>et al.</i>	2016	India	15	Cord	23.8 (22.2-25.4)	
Basu <i>et al.</i>	2015	India	142	Cord	26.5 (25.5-27.5)	
Bermudez <i>et al.</i>	2015	Spain	30	Cord	6.26 (5.37-7.15)	
Briana <i>et al.</i>	2013	Greece	104	Cord	24.14 (22.4-25.9)	
Busarira <i>et al.</i>	2019	Libya	126	Cord	23.69 (23.5-23.9)	
Cao <i>et al.</i>	2016	USA	68	Cord	39.73 (35-44.4)	
Chong <i>et al.</i>	1984	UK	20	Cord	41.1 (29.6-52.6)	
Delaney <i>et al.</i>	2019	USA	108	Cord	40.8 (37.3-44.3)	
El-Farrash <i>et al.</i>	2012	Egypt	30	Cord	28.29 (25.6-31)	
Ertekin <i>et al.</i>	2015	Turkey	76	Cord	26.1 (24.1-28.1)	
Ervasti <i>et al.</i>	2007	Finland	199	Cord	27.4 (26.3-28.5)	
Gruccio <i>et al.</i>	2014	Argentina	99	Cord	27.03 (25.7-28.4)	
Haga <i>et al.</i>	1980	Norway	21	Cord	27.1 (24.2-30)	
Kelly <i>et al.</i>	1978	Scotland	115	Cord	27.0 (25.6-28.4)	

Kleven <i>et al.</i>	2007	USA	26	Cord	44.1 (32.2-56)	
Kocylowski <i>et al.</i>	2017	Poland	64	Cord	35.1 (33.2-37)	
Lao <i>et al.</i>	1991	Hong Kong	77	Cord	35.8 (32.7--38.9)	
Lee <i>et al.</i>	2006	South Korea	19	Cord	31.3 (28.4-34.2)	
Lee <i>et al.</i>	2016	USA	104	Cord	35.4 (32-39.2)	
Mezdoud <i>et al.</i>	2017	Algeria	97	Cord	20.1 (19-21.3)	
Milman <i>et al.</i>	1987	Denmark	74	Cord		28 (19-39)
			47	Venous - 120 hours		19 (13-31)
Mukhopadhyay <i>et al.</i>	2011	India	50	Cord	29 (25.8-32.2)	
Murata <i>et al.</i>	1989	Japan	45	Cord	28.5 (26.7-30.3)	
Oliveria <i>et al.</i>	2014	Brazil	144	Cord	24.6 (23.5-25.7)	
Ozkiraz <i>et al.</i>	2011	Turkey	16	Venous - 216 hours (96-336)	16.4 (13.8-19)	
Patidar <i>et al.</i>	2013	India	50	Venous - 8 hours (1-23)	19.4 (17.2-21.6)	
Prentice <i>et al.</i>	2019	The Gambia	81	Cord	24.7 (22.5-26.9)	
			53	Venous - 6 hours (2-11)	13.6 (12.0-15.2)	
			21	Venous - 29 hours (26-34)	11.6 (10.1-13.1)	
			33	Venous - 77 hours (74-82)	14.5 (13.1-16.0)	
Puolakka <i>et al.</i>	1980	Finland	47	Cord	28.8 (26.2-31.4)	
Rois <i>et al.</i>	1975	USA	26	Cord	6.19 (6.18-6.20)	
Ru <i>et al.</i>	2018	USA	49	Cord	48.3 (39.3-59.1)	
Slomka <i>et al.</i>	2013	Poland	49	Cord	22.4 (20.5-24.3)	
Sweet <i>et al.</i>	2001	UK	68	Cord	26 (24.2-27.8)	
Szabo <i>et al.</i>	2001	Hungary	10	Cord	23.2 (16.3-30.1)	
			10	Venous - 47 hours	7.2 (6.15-8.25)	
Tiker <i>et al.</i>	2006	Turkey	16	Venous - 209 hours (96-288)	19.9 (12.7-28.4)	
Tsuzuki <i>et al.</i>	2013	Japan	30	Cord	31.1 (27.3-34.9)	
			30	Venous - 120 hours	19.5 (16.4-22.6)	
Yamada <i>et al.</i>	2014	Brazil	21	Cord	23.9 (19.4-28.4)	

			21	Venous - 720 hours	16.7 (14.9-18.5)	
Yapakci <i>et al.</i>	2009	Turkey	16	Venous - 211 hours (±46)	19.9 (17.9-21.9)	
<b>Weighted Mean (Cord)</b>					<b>28.5 (26.0-31.1)</b>	<b>N/A</b>
<b>Weighted Mean (Venous)</b>					<b>13.8 (10.8-16.9)</b>	<b>N/A</b>
<b>Adults (Median (95% CI))</b>					<b>15.2 (8.1-24.5) µmol/L (10-90th percentiles) - NHNES</b>	
<b>Infants (Median (95% CI))</b>					<b>12.5 (5.5-20.6) µmol/L (10-90th percentiles) - NHNES</b>	

**Table 7: Serum iron concentration (µmol/L) in preterm newborns over the neonatal period.**

Reference	Year	Location	n	Type of Sample (Cord or Venous)	Study Group	Serum Iron (µmol/L)	
						Mean (95% CI or Range)	Median (IQR or 95% CI)
Celik <i>et al.</i>	2015	Turkey	42	Venous - 648 hours (288-1872)		15.6 (13.3-17.9)	
Delaney <i>et al.</i>	2019	USA	126	Cord		53.1 (49.8-56.4)	
Haga <i>et al.</i>	1980	Norway	24	Cord	AGA Group <sup>c</sup>	16.8 (13.2-20.4)	
			7	Cord	SGA Group <sup>d</sup>	18.3 (11.2-25.4)	
Ichinomiya <i>et al.</i>	2017	Japan	92	Cord			23.27 (15.2-32.4)
Lackmann <i>et al.</i>	1998	Germany	15	Venous - <1 hour	<32 wks		14 (2-41)
			22	Venous - <1 hour	33-34 wks		12 (2-32)
			26	Venous - <1 hour	35-36 wks		15 (6-28)
Ru <i>et al.</i>	2018	USA	91	Cord	<sup>e</sup>	73.4 (57.3-93.1)	
Ru <i>et al.</i>	2018	USA	140	Cord	<sup>f</sup>	53.7 (50.1-60.8)	
Schiza <i>et al.</i>	2007	Greece	181	Venous - 336 hours		16.1 (15.5-16.7)	
Sweet <i>et al.</i>	2001	UK	50	Cord	30-36 wks <sup>g</sup>	20.8 (18.4-23.2)	
			26	Cord	24-29 wks <sup>h</sup>	17.4 (13.3-21.5)	
Tiker <i>et al.</i>	2006	Turkey	14	Venous - 67 hours (24-144)	26-32 wks	15.81 (4.83-33.48)	
			12	Venous - 65 hours (24-96)	33-36 wks	19.26 (6.8-39.2)	
Tsuzuki <i>et al.</i>	2013	Japan	14	Cord		27.5 (21.9-33.1)	
				Venous - 120 hours		16.47 (13.1-19.8)	
Yamada <i>et al.</i>	2014	Brazil	25	Cord		8.8 (6.97-10.6)	
				Venous - 720 hours		15.54 (14.1-17)	
Yapakci <i>et al.</i>	2009	Turkey	17	Venous - 336 hours (96-720)		17.22 (14.2-20.3)	
<b>Weighted Mean (Cord)</b>						<b>46.8 (29.7-63.8)</b>	<b>N/A</b>
<b>Weighted Mean (Venous)</b>						<b>16.2 (15.3-17.0)</b>	<b>N/A</b>

Adults (Median (95% CI))	15.2 (8.1-24.5) $\mu\text{mol/L}$ (10-90th percentiles) - NHNES
Infants (Median (95% CI))	12.5 (5.5-20.6) $\mu\text{mol/L}$ (10-90th percentiles) - NHNES

## FIGURE LEGENDS

**Figure 1: Placental iron transfer between mother and fetus.**  $\text{Fe}^{2+}$  = ferrous iron,  $\text{Fe}^{3+}$  = ferric iron, Tf = transferrin, Apo-Tf = unsaturated transferrin, Fetal Tf = fetal-derived transferrin, NTBI = non-transferrin bound iron. Syncytiotrophoblasts in the placental villi take up transferrin-bound iron from the maternal circulation by endocytosis via transferrin receptor 1 (TFR1). Iron is released from TFR1 in acidified endosomes and transferred into the syncytiotrophoblast cytoplasm by DMT-1, Zrt/Irt-like protein ZIP8, and ZIP14, collectively. Ferroportin transports iron out of placental syncytiotrophoblasts, and then ceruloplasmin, hephaestin, and zyklopen oxidise  $\text{Fe}^{2+}$  to  $\text{Fe}^{3+}$  helping it pass through the endothelium to reach the fetal circulation. It is still unclear as to whether newly transported iron enters the fetal circulation as NTBI or bound to fetal transferrin. Fetal-derived hepcidin is believed to regulate ferroportin expression on the fetal basal-side of placental syncytiotrophoblasts.<sup>13,117</sup> Maternal-derived hepcidin is believed to play a role in regulating TFR1 expression on the maternal-side of the placental syncytiotrophoblasts<sup>118</sup>. This figure is adapted from Sangkhae and Nemeth *et al*, 2018.<sup>119</sup>

**Figure 2: Flow diagram of the literature search and selection criteria.** Retrieving publications on hepcidin, TSAT or serum iron in neonates over the first month of life.

**Figure 3: Hepcidin (ng/mL) in cord blood:** (A) full term neonates, (B) preterm neonates. Standardized means from each publication are plotted with error bars showing 95% confidence intervals. Dashed line shows the weighted mean of all publications found in the figure. Median values were not included in this figure.



**Figure 4: Transferrin saturation (%) in cord blood:** (A) full term neonates, (B) preterm neonates. Means from each publication are plotted with error bars showing 95% confidence intervals. Dashed line shows the weighted mean of all publications found in the figure. <sup>a</sup> shows Haga *et al*, AGA group.<sup>120</sup> <sup>b</sup> shows Haga *et al* SGA group.<sup>120</sup> Median values were not included in this figure.

**Figure 5: Serum iron (μmol/L) in cord blood:** (A) full term neonates, (B) preterm neonates. Means from each publication are plotted with error bars showing 95% confidence intervals. Dashed line shows the weighted mean of all publications found in the figure. <sup>c</sup> shows AGA neonates in Haga *et al*.<sup>120</sup> <sup>d</sup> shows SGA neonates in Haga *et al*.<sup>120</sup> Ru *et al*, 2018<sup>115</sup> is referenced as <sup>e</sup>. Ru *et al*, 2018<sup>116</sup> is referenced as <sup>f</sup>. <sup>g</sup> shows 30-36 wks neonates in Sweet *et al*.<sup>121</sup> <sup>h</sup> shows 24-29 wks neonates in Sweet *et al*.<sup>121</sup> Median values were not included in this figure.

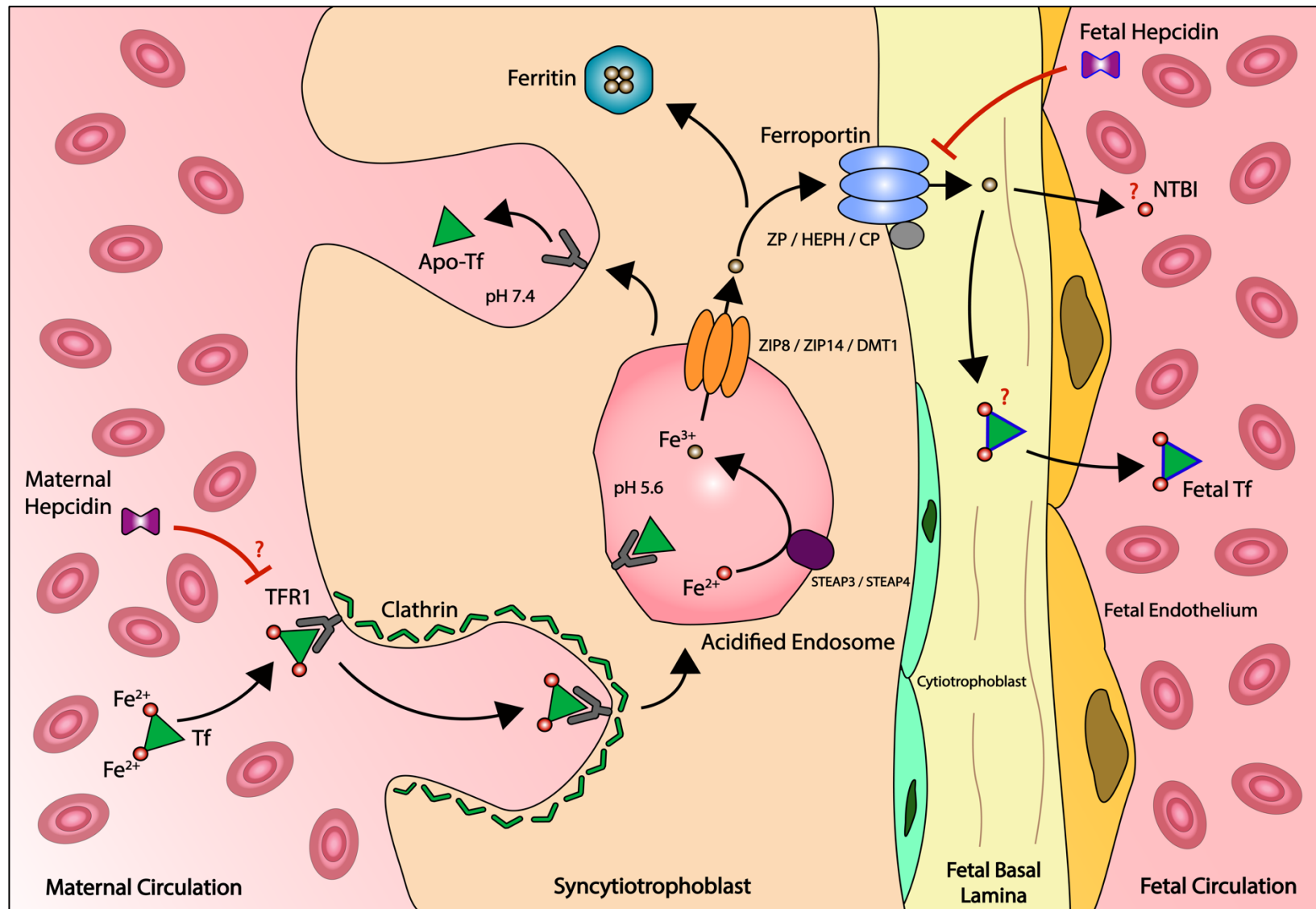
**Figure 6: Hepcidin (ng/mL) over the neonatal period:** (A) full term neonates,  $\alpha$  shows the weighted mean (95%CI) for all studies seen in **Figure 2A**.  $\beta$ ,  $\chi$  and  $\varepsilon$  shows Prentice *et al*.<sup>108</sup>  $\delta$  shows Kulik-Rechberger *et al*.<sup>26</sup> (B) preterm neonates,  $\alpha$  shows the weighted mean (95%CI) for all studies seen in Figure 2B.  $\beta$  shows Uijterschout *et al*.<sup>122</sup>

**Figure 7: Transferrin saturation (%) over the neonatal period:** (A) full term neonates,  $\alpha$  shows the weighted mean (95%CI) for all studies seen in **Figure 3A**.  $\beta$  shows Prentice *et al*.<sup>108</sup>  $\chi$  shows Al-Tawil *et al*.<sup>123</sup>  $\delta$  shows Prentice *et al*.<sup>108</sup>  $\varepsilon$  shows Balogh *et al*.<sup>101</sup>  $\phi$  shows Anderson *et al*.<sup>124</sup>  $\gamma$  shows Prentice *et al*.<sup>108</sup>  $\eta$  shows Milman *et al*.<sup>40</sup>  $\iota$  shows Kitajima *et al*.<sup>125</sup>  $\varphi$  shows Yamada *et al*.<sup>126</sup> (B) preterm neonates,  $\alpha$  shows the

weighted mean (95%CI) for all studies seen in **Figure 3B**.  $\beta$  shows Lackmann *et al.*<sup>127</sup>  $\chi$  shows Celik *et al.*<sup>128</sup>  $\delta$  shows Yamada *et al.*<sup>126</sup>  $\varepsilon$  shows Kitajima *et al.*<sup>125</sup> All values are mean (95%CI), unless marked with  $\circ$  median (range) and  $\bullet$  median (95%CI). Lackmann *et al.*, 1998 ( $\beta$ ) data from the three study groups (<32 wks, 33-34 wks and 35-36 wks) was averaged as all groups are classed as PTB neonates and are bled at the same time of life.<sup>127</sup>

**Figure 8: Serum iron ( $\mu\text{mol/L}$ ) over the neonatal period:** (A) full term neonates,  $\alpha$  shows the weighted mean (95%CI) for all studies seen in **Figure 2A**.  $\beta$  shows Prentice *et al.*<sup>108</sup>  $\chi$  shows Patidar *et al.*<sup>129</sup>  $\delta$  shows Prentice *et al.*<sup>108</sup>  $\varepsilon$  shows Balogh *et al.*<sup>101</sup>  $\phi$  shows Szabo *et al.*<sup>130</sup>  $\gamma$  shows Anderson *et al.*<sup>124</sup>  $\eta$  shows Prentice *et al.*<sup>108</sup>  $\iota$  shows Milman *et al.*<sup>40</sup>  $\varphi$  shows Tsuzuki *et al.*<sup>131</sup>  $\kappa$  shows Tiker *et al.*<sup>132</sup>  $\lambda$  shows Yapakci *et al.*<sup>133</sup>  $\mu$  shows Ozkiraz *et al.*<sup>134</sup>  $\nu$  shows Yamada *et al.*<sup>126</sup> (B) preterm neonates,  $\alpha$  shows the weighted mean (95%CI) for all studies seen in **Figure 2B**.  $\beta$  shows Lackmann *et al.*<sup>127</sup>  $\chi$  shows Tiker *et al.*<sup>132</sup>  $\delta$  shows Tiker *et al.*<sup>132</sup>  $\varepsilon$  shows Tsuzuki *et al.*<sup>131</sup>  $\phi$  shows Schiza *et al.*<sup>135</sup>  $\gamma$  shows Yapakci *et al.*<sup>133</sup>  $\eta$  shows Celik *et al.*<sup>128</sup>  $\iota$  shows Yamada *et al.*<sup>126</sup> All values are mean (95%CI), unless marked with \* mean (range),  $\circ$  median (range) and  $\bullet$  median (95%CI). Lackmann *et al.*, 1998 ( $\beta$ ) data from the three study groups (<32 wks, 33-34 wks and 35-36 wks) was averaged as all groups are classed as PTB neonates and are bled at the same time of life.<sup>127</sup>

Figure 1: Placental iron transfer between mother and fetus



**Figure 2: Flow diagram of the literature search.**

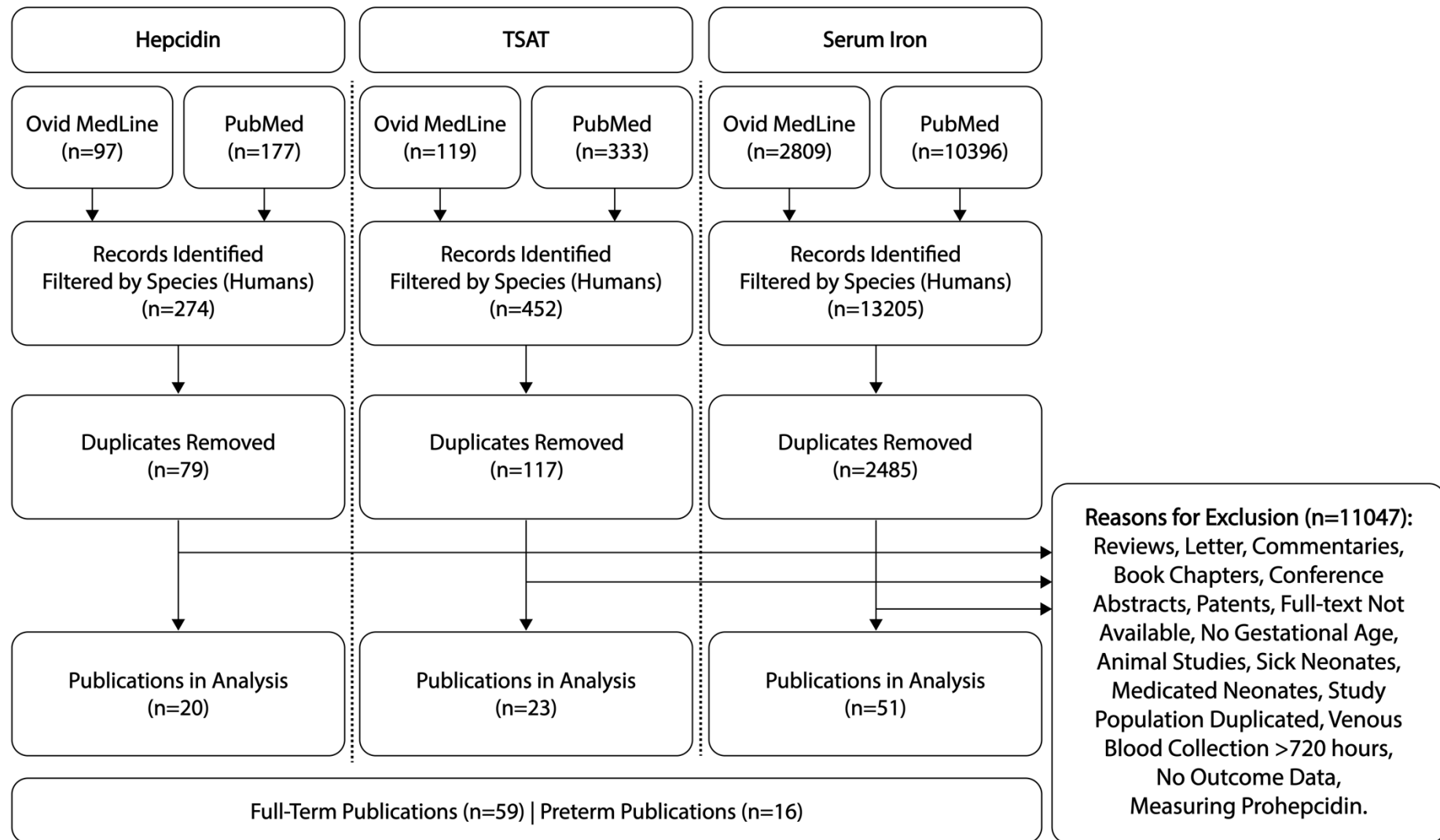


Figure 3: Standardized hepcidin concentration (ng/mL) in cord blood.

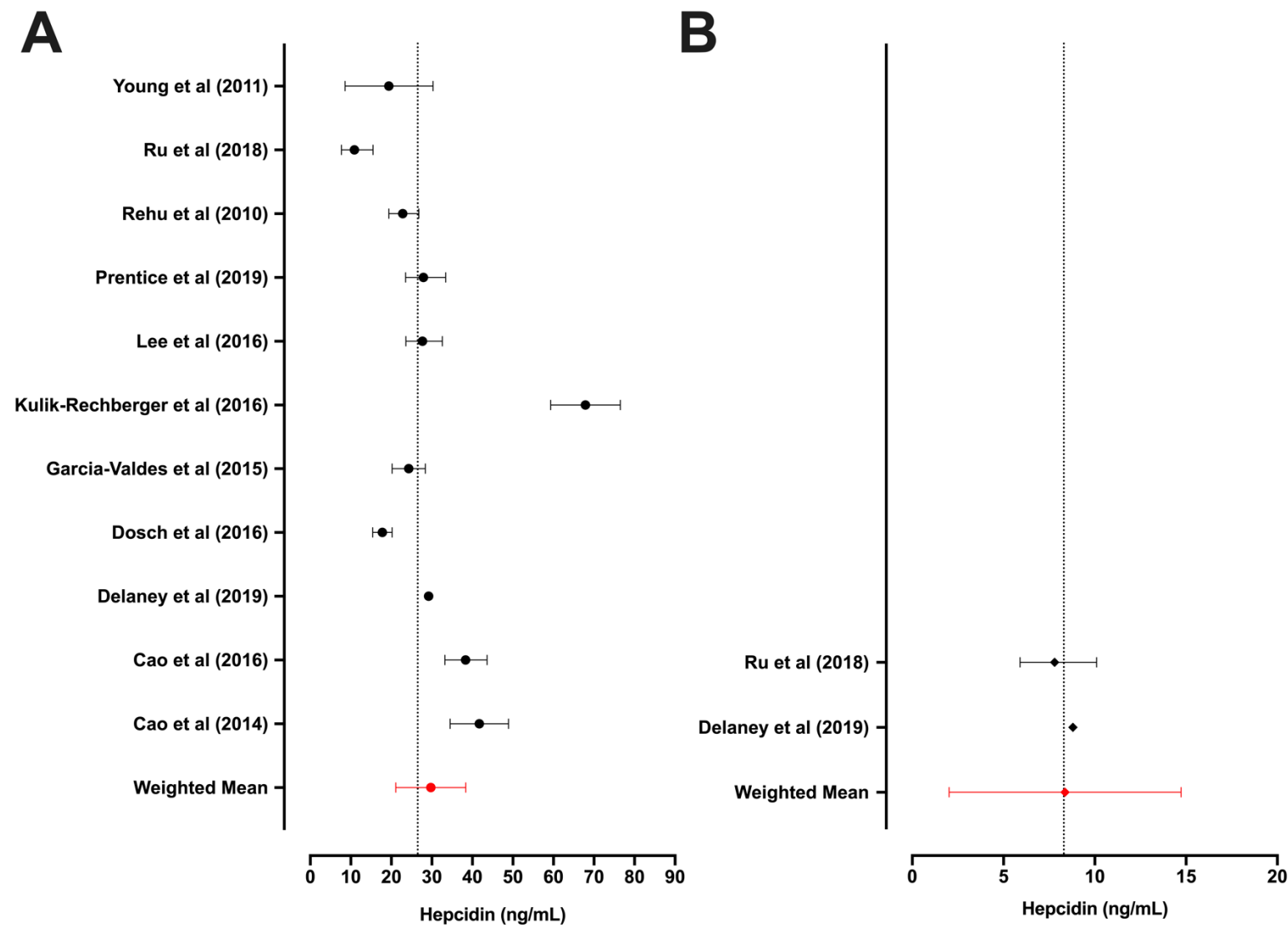


Figure 4: Transferrin saturation (%) in cord blood.

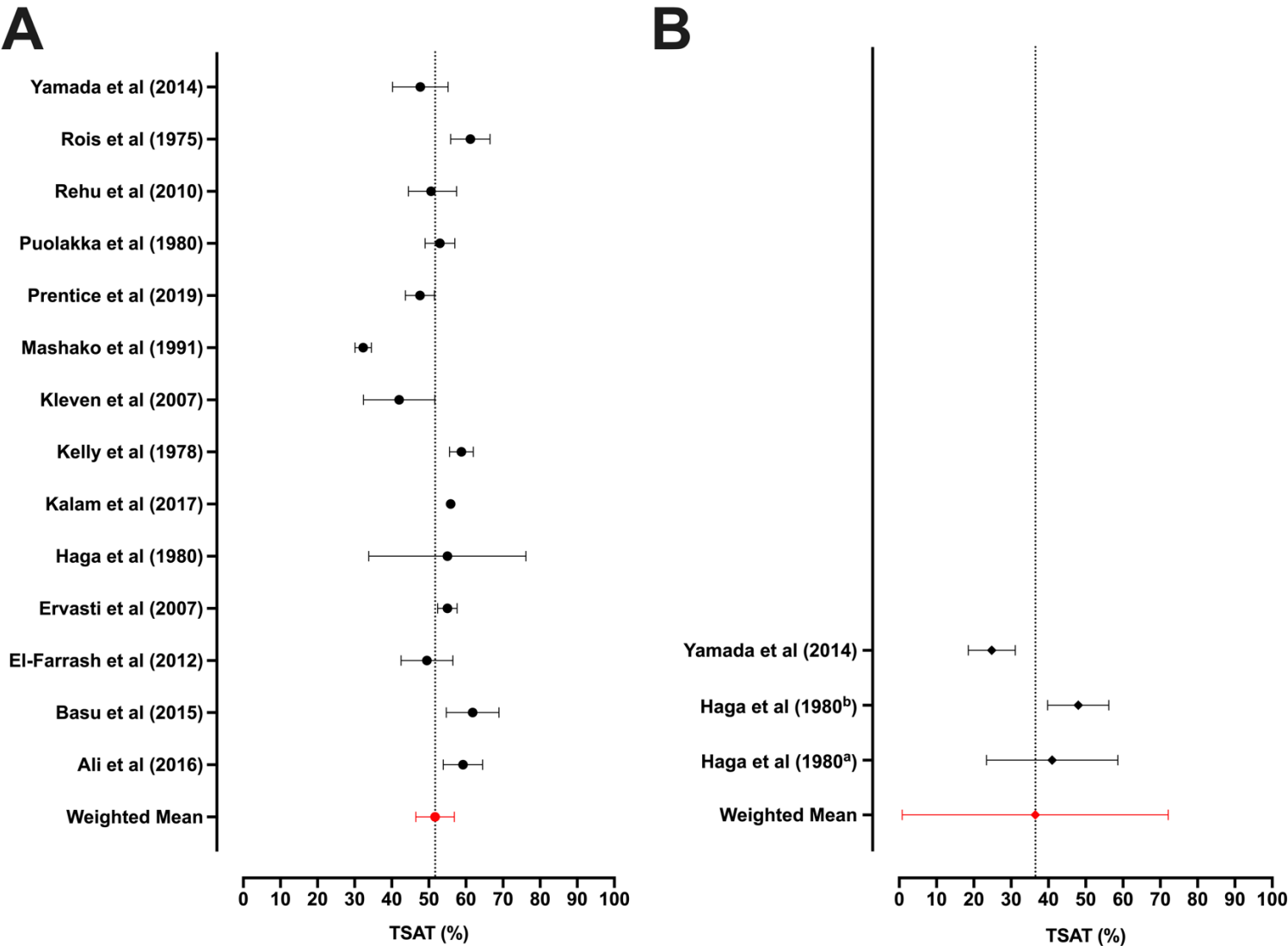


Figure 5: Serum iron concentration ( $\mu\text{mol/L}$ ) in cord blood

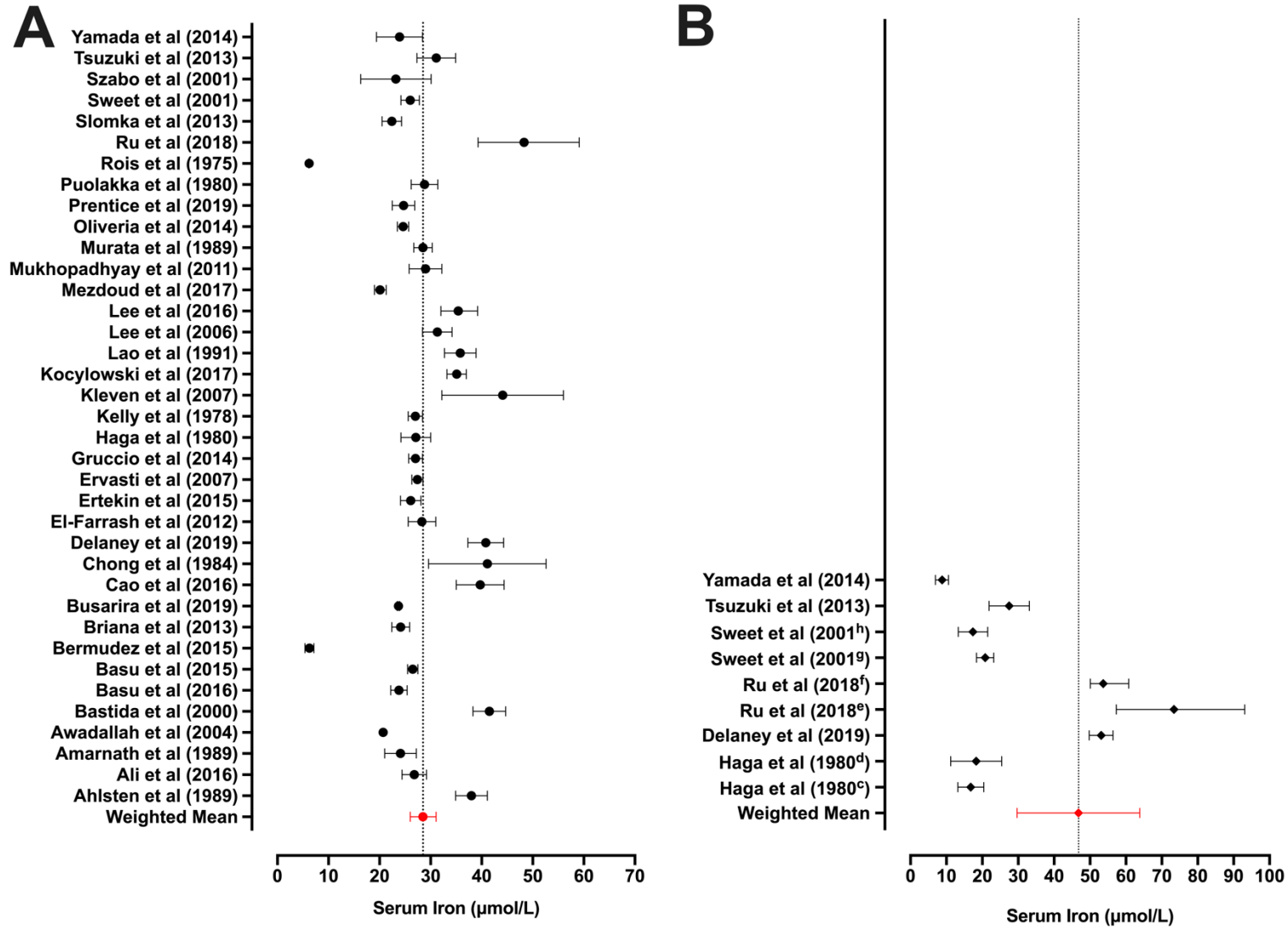


Figure 6: Hepcidin concentration (ng/mL) over the neonatal period

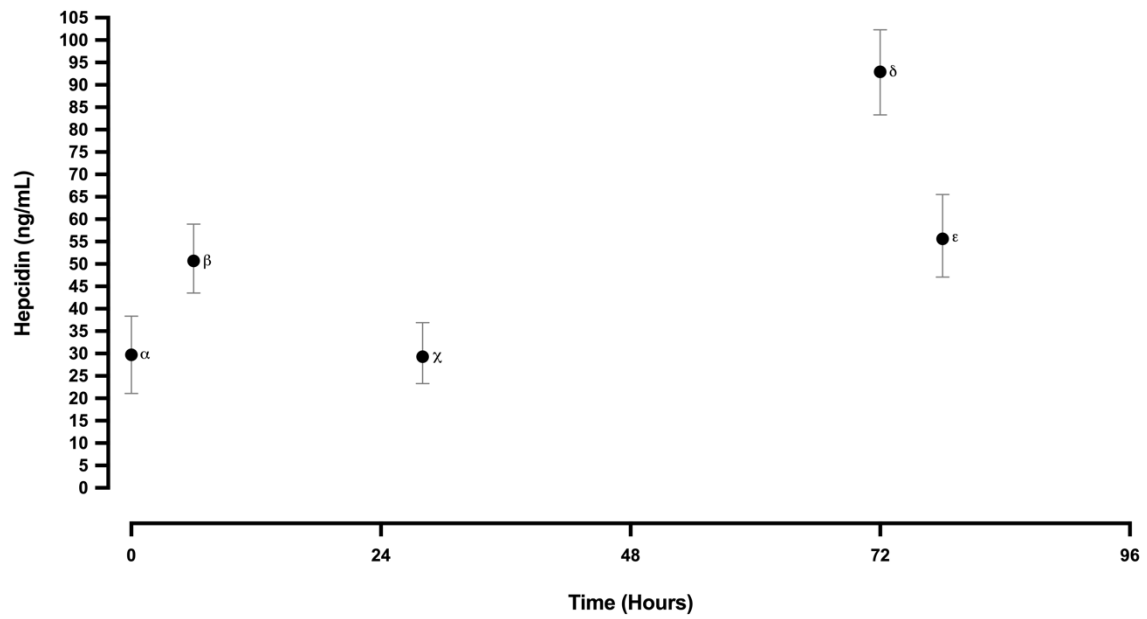


Figure 6A - Hepcidin in FTB neonates

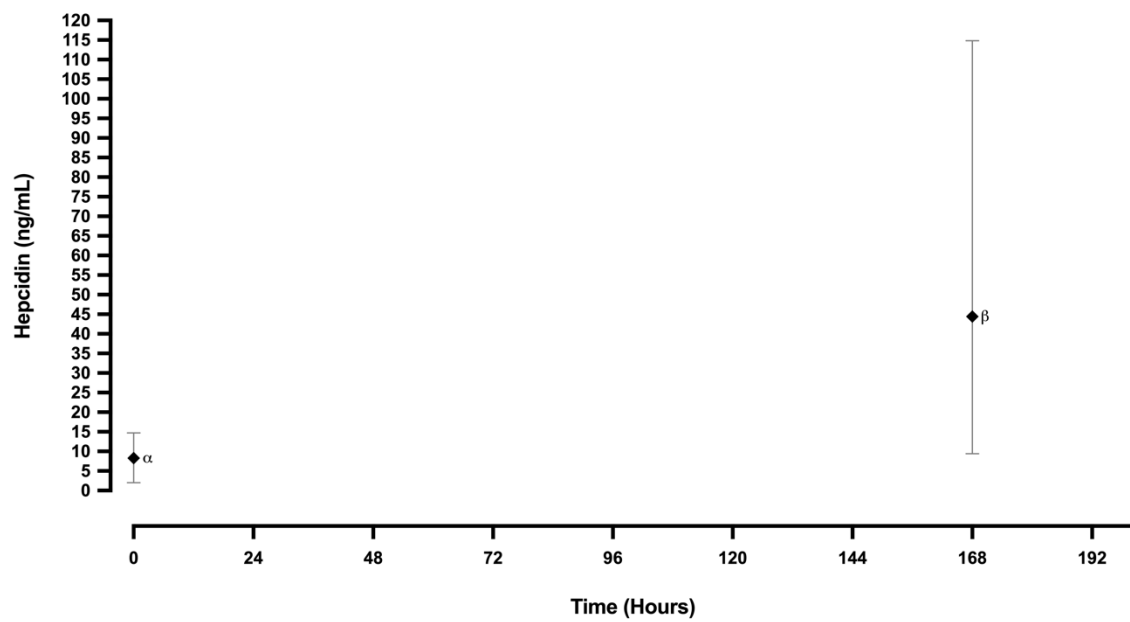
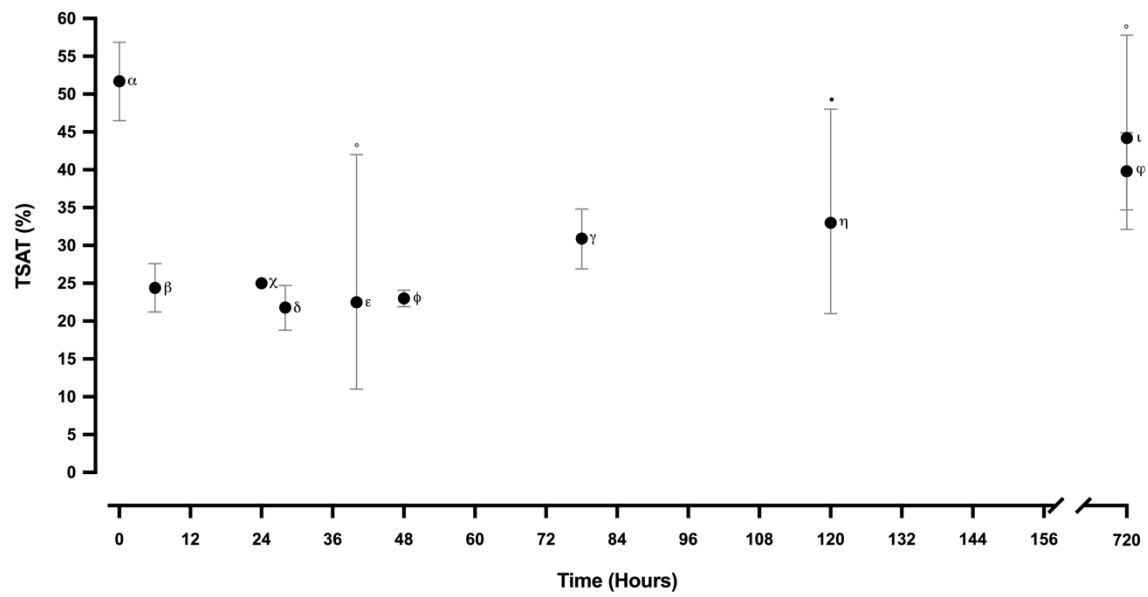


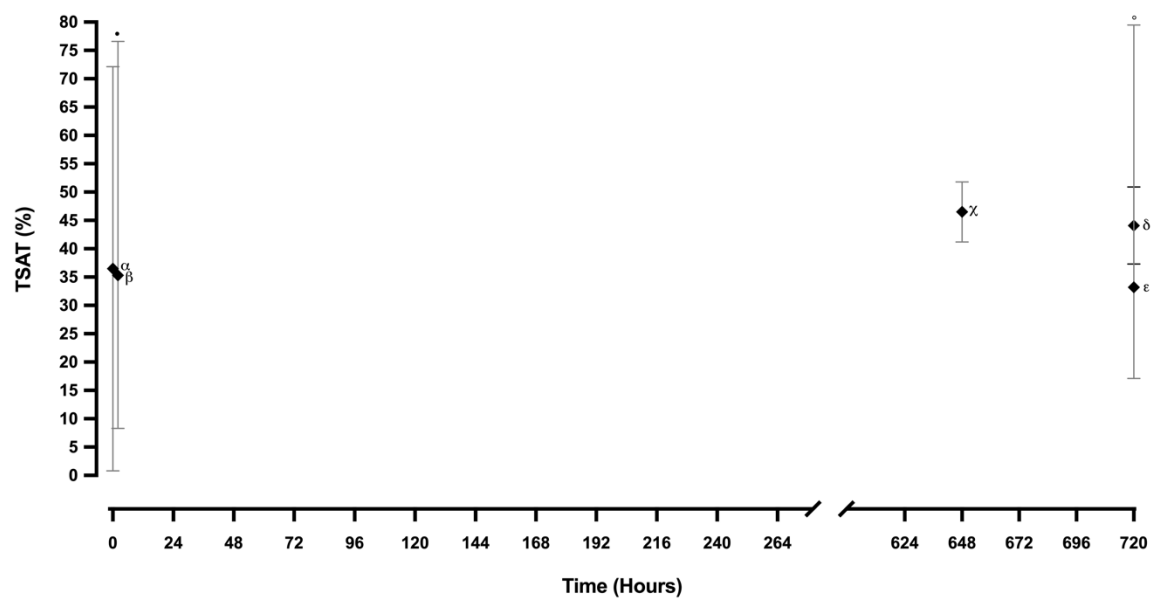
Figure 6B - Hepcidin in PTB neonates



**Figure 7: Transferrin saturation (%) over the neonatal period**

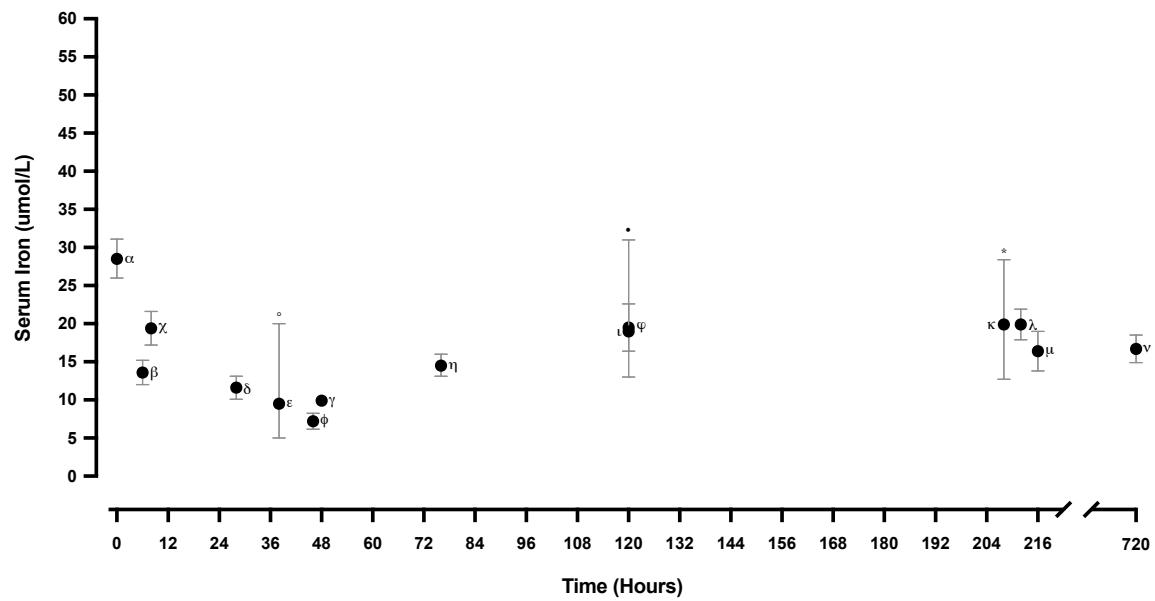


**Figure 7A - TSAT in FTB neonates**

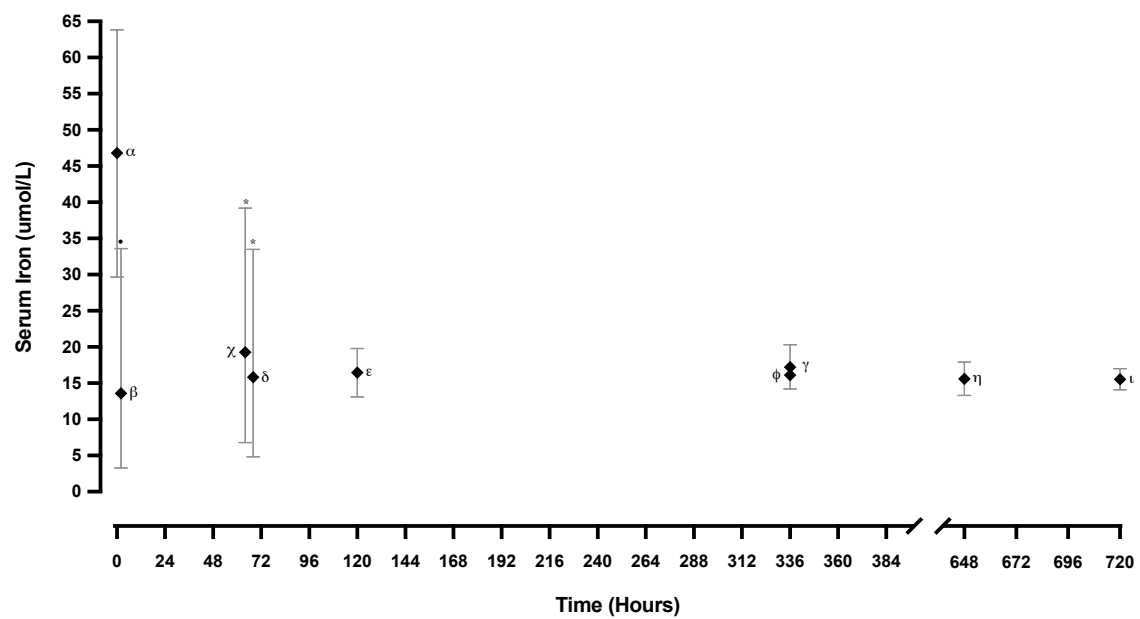


**Figure 7B - TSAT in PTB neonates**

**Figure 8: Serum iron concentration ( $\mu\text{mol/L}$ ) over the neonatal period**



**Figure 8A - Serum iron in FTB neonates**



**Figure 8B - Serum iron in PTB neonates**

## Title

**Hepcidin, serum iron and transferrin saturation in full term and premature infants during the first month of life: A review of existing evidence in humans.**

## Authors

James H. Cross, BSc<sup>1</sup>, Andrew M. Prentice, PhD<sup>1</sup>, Carla Cerami, MD<sup>1\*</sup>.

## Affiliations

<sup>1</sup>Medical Research Council Unit The Gambia at the London School of Hygiene & Tropical Medicine, Atlantic Boulevard, Fajara, P.O. Box 273, Banjul, The Gambia.

## 3.7 SUPPLEMENTARY MATERIAL

### SUPPLEMENTARY METHODS

#### Supplementary Table 1 - Post-hoc hepcidin standardisation

Test ID	Kit	Slope (95% CI)	Intercept (95% CI)
IC-4	Bachem Hepcidin-25	1.569 (1.523-1.614)	-0.10 (-1.95 to 1.85)
IC-5	Intrinsic Lifesciences Hepcidin IDx	3.184 (3.086-3.281)	-0.85 (-2.91 to 1.21)
IC-6	DRG Hepcidin-25	0.711 (0.674-0.748)	0.73 (-0.05 to 1.52)

**Supplementary Table 1:** Previously documented regression relationships used to conduct post-hoc hepcidin standardisation. These regression relationships were extract from **Table 2** of van der Vorm *et al.*<sup>1</sup>

### SUPPLEMENTARY REFERENCES

1. van der Vorm LN, Hendriks JCM, Laarakkers CM, Klaver S, *et al.* Toward Worldwide Hepcidin Assay Harmonization: Identification of a Commutable Secondary Reference Material. *Clin Chem. Clinical Chemistry*; 2016;62:993–1001.

## **Chapter 4 - Neonatal iron distribution and infection susceptibility in full term, preterm and low birthweight babies in urban Gambia: study protocol for an observational study (Methodology)**

---

### **Summary of Chapter**

**BACKGROUND:** Neonatal infection is the third largest cause of death in children under-five worldwide. Nutritional immunity is the process by which the host innate immune system limits nutrient availability to invading organisms. Iron is an essential micronutrient for both microbial pathogens and their mammalian hosts. Changes in iron availability and distribution have significant effects on pathogen virulence and on the immune response to infection. Our previously published data shows that, during the first 24 hours of life, full-term neonates have reduced overall serum iron. Transferrin saturation decreases rapidly from 45% in cord blood to ~20% by six hours post-delivery.

**METHODS:** To study neonatal nutritional immunity and its role in neonatal susceptibility to infection, we will conduct an observational study on 300 full-term normal birthweight (FTB+NBW), 50 preterm normal birthweight (PTB+NBW), 50 preterm low birthweight (PTB+LBW) and 50 full-term low birthweight (FTB+LBW), vaginally-delivered neonates born at Kanifing General Hospital, The Gambia. We will characterize and quantify iron-related nutritional immunity during the early neonatal period and use *ex vivo* sentinel bacterial growth assays to assess how differences in serum iron affect bacterial growth. Blood samples will be collected from the umbilical cord (arterial and venous) and at serial time points from the neonates over the first week of life.

**DISCUSSION:** Currently, little is known about nutritional immunity in neonates. In this study, we will increase understanding of how nutritional immunity may protect neonates from

infection during the first critical days of life by limiting the pathogenicity and virulence of neonatal sepsis causing organisms by reducing the availability of iron. Additionally, we will investigate the hypothesis that this protective mechanism may not be activated in preterm and low birthweight neonates, potentially putting these babies at an enhanced risk of neonatal infection.

**\*NOTE:** The secondary study objective (III) of assessing the bacterial growth of common neonatal pathogens in neonatal sera was not achieved due to the lack of sufficient blood sample volume.

## RESEARCH PAPER COVER SHEET - METHODOLOGY PAPER

---

Please note that a cover sheet must be completed for each research paper included within a thesis.

### **SECTION A – Student Details**

<b>Student ID Number</b>	LSH158152	<b>Title</b>	Mr.
<b>First Name(s)</b>	James Henry		
<b>Surname/Family Name</b>	Cross		
<b>Thesis Title</b>	Iron and Infection: Neonatal Iron Transition		
<b>Primary Supervisor</b>	Dr Carla Cerami		

If the Research Paper has previously been published please complete Section B, if not please move to Section C.

### **SECTION B – Paper already published**

Where was the work published?	Gates Open Research		
When was the work published?	21 <sup>st</sup> May 2019 (First Published) 15 <sup>th</sup> October 2019 (Latest Published)		
If the work was published prior to registration for your research degree, give a brief rationale for its inclusion	N/A		
Have you retained the copyright for the work?*	<b>Yes</b>	Was the work subject to academic peer review?	<b>Yes</b>

\*If yes, please attach evidence of retention. If no, or if the work is being included in its published format, please attach evidence of permission from the copyright holder (publisher or other author) to include this work.

**Keywords**

Nutritional Immunity, Host-Pathogen Interaction, Hepcidin, Neonates, Hypoferremia, Transferrin, The Gambia, Sub-Saharan Africa

**Corresponding author:** Carla Cerami ([ccerami@mrc.gm](mailto:ccerami@mrc.gm))

**Author roles:** **Cross JH:** Data Curation, Formal Analysis, Investigation, Project Administration, Writing – Original Draft Preparation, Writing – Review & Editing; **Jarjou O:** Investigation, Project Administration, Supervision, Writing – Review & Editing; **Mohammed NI:** Formal Analysis, Writing – Review & Editing; **Prentice AM:** Conceptualization, Formal Analysis, Writing – Review & Editing; **Cerami C:** Conceptualization, Data Curation, Formal Analysis, Funding Acquisition, Investigation, Methodology, Project Administration, Resources, Supervision, Writing – Original Draft Preparation, Writing – Review & Editing

**Competing interests:** No competing interests were disclosed.

**Grant information:** This study was supported by the Bill & Melinda Gates Foundation [OPP1152353].

*The funders had no role in study design, data collection and analysis, decision to publish, or preparation of the manuscript.*

**Copyright:** © 2019 Cross JH *et al.* This is an open access article distributed under the terms of the [Creative Commons Attribution Licence](#), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

**How to cite this article:** Cross JH, Jarjou O, Mohammed NI *et al.* Neonatal iron distribution and infection susceptibility in full term, preterm and low birthweight babies in urban Gambia: study protocol for an observational study. [version 1; peer review: awaiting peer review] Gates Open Research 2019, 3:1469 (<https://doi.org/10.12688/gatesopenres.12963.1>)

**First published:** 21 May 2019, 3:1469 (<https://doi.org/10.12688/gatesopenres.12963.1>)

**SECTION C – Prepared for publication, but not yet published**

Where is the work intended to be published?

Please list the paper's authors in the intended authorship order:

Stage of publication

Choose an item.

**SECTION D – Multi-authored work**

For multi-authored work, give full details of your role in the research included in the paper and in the preparation of the paper. (Attach a further sheet if necessary)

This paper was proposed at the start of the study by myself and my supervisor to reference all materials and methods for the subsequent original publications. I coordinated the development of study protocols and data analysis plan. Additionally, I produced the first draft of this paper for review by co-authors. Dr Carla Cerami, Dr Ousman Jarjou and Professor Andrew Prentice all helped re-structure the first draft and provided suggestions towards making edits. Dr Nuredin Ibrahim Mohammed advised in the formal statistical analysis plan.

**SECTION E**

Student Signature



Date

08/08/2019

**Supervisor Signature**



**Date**

08/08/2019





## STUDY PROTOCOL

# Neonatal iron distribution and infection susceptibility in full term, preterm and low birthweight babies in urban Gambia: study protocol for an observational study. [version 1; peer review: 2 approved]

James H. Cross, Ousman Jarjou , Nuredin Ibrahim Mohammed ,  
Andrew M. Prentice, Carla Cerami

MRC Unit The Gambia at the London School of Hygiene & Tropical Medicine, Fajara, The Gambia

**v1** First published: 21 May 2019, 3:1469 (<https://doi.org/10.12688/gatesopenres.12963.1>)

Latest published: 15 Oct 2019, 3:1469 (<https://doi.org/10.12688/gatesopenres.12963.2>)

## Abstract

**Background:** Neonatal infection is the third largest cause of death in children under five worldwide. Nutritional immunity is the process by which the host innate immune system limits nutrient availability to invading organisms. Iron is an essential micronutrient for both microbial pathogens and their mammalian hosts. Changes in iron availability and distribution have significant effects on pathogen virulence and on the immune response to infection. Our previously published data shows that, during the first 24 hours of life, full-term neonates have reduced overall serum iron. Transferrin saturation decreases rapidly from 45% in cord blood to ~20% by six hours post-delivery.

**Methods:** To study neonatal nutritional immunity and its role in neonatal susceptibility to infection, we will conduct an observational study on 300 full-term normal birth weight (FTB+NBW), 50 preterm normal birth weight (PTB+NBW), 50 preterm low birth weight (PTB+LBW) and 50 full-term low birth weight (FTB+LBW), vaginally-delivered neonates born at Kanifing General Hospital, The Gambia. We will characterize and quantify iron-related nutritional immunity during the early neonatal period and use *ex vivo* sentinel bacterial growth assays to assess how differences in serum iron affect bacterial growth. Blood samples will be collected from the umbilical cord (arterial and venous) and at serial time points from the neonates over the first week of life.

**Discussion:** Currently, little is known about nutritional immunity in neonates. In this study, we will increase understanding of how nutritional immunity may protect neonates from infection during the first critical days of life by limiting the pathogenicity and virulence of neonatal sepsis causing organisms by reducing the availability of iron. Additionally, we will investigate the hypothesis that this protective mechanism may not be activated in preterm and low birth weight neonates, potentially putting these babies at an enhanced risk of neonatal infection.

**Trial registration:** clinicaltrials.gov (NCT03353051) 27/11/2017

## Open Peer Review

Reviewer Status

Invited Reviewers

1 2

REVISED

### version 2

published  
15 Oct 2019

### version 1

published  
21 May 2019

report

report

1 **Deborah Stefanova**, ETH Zurich, Zurich, Switzerland

2 **Veena Sangkhae** , University of California, Los Angeles (UCLA), Los Angeles, USA

Any reports and responses or comments on the article can be found at the end of the article.

## Keywords

Nutritional Immunity, Host-Pathogen Interaction, Hepcidin, Neonates, Hypoferremia, Transferrin, The Gambia, Sub-Saharan Africa

**Corresponding author:** Carla Cerami ([ccerami@mrc.gm](mailto:ccerami@mrc.gm))

**Author roles:** **Cross JH:** Data Curation, Formal Analysis, Investigation, Project Administration, Writing – Original Draft Preparation, Writing – Review & Editing; **Jarjou O:** Investigation, Project Administration, Supervision, Writing – Review & Editing; **Mohammed NI:** Formal Analysis, Writing – Review & Editing; **Prentice AM:** Conceptualization, Formal Analysis, Writing – Review & Editing; **Cerami C:** Conceptualization, Data Curation, Formal Analysis, Funding Acquisition, Investigation, Methodology, Project Administration, Resources, Supervision, Writing – Original Draft Preparation, Writing – Review & Editing

**Competing interests:** No competing interests were disclosed.

**Grant information:** This study was supported by the Bill & Melinda Gates Foundation [OPP1152353].

*The funders had no role in study design, data collection and analysis, decision to publish, or preparation of the manuscript.*

**Copyright:** © 2019 Cross JH *et al.* This is an open access article distributed under the terms of the [Creative Commons Attribution License](#), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

**How to cite this article:** Cross JH, Jarjou O, Mohammed NI *et al.* Neonatal iron distribution and infection susceptibility in full term, preterm and low birthweight babies in urban Gambia: study protocol for an observational study. [version 1; peer review: 2 approved] Gates Open Research 2019, 3:1469 (<https://doi.org/10.12688/gatesopenres.12963.1>)

**First published:** 21 May 2019, 3:1469 (<https://doi.org/10.12688/gatesopenres.12963.1>)

## Abbreviations

PTB = Preterm birth  
 FTB = Fullterm birth  
 LBW = Low birthweight  
 NBW = Normal birthweight  
 WHO = World Health Organisation  
 TSAT = Transferrin saturation  
 MDG = Millennium development goal  
 GBS = Group B *Streptococci*  
 EONS = Early onset neonatal sepsis  
 YICSS = Young Infants Clinical Signs Studies  
 KGH = Kanifing General Hospital  
 MRCG = Medical Research Council Unit The Gambia at LSHTM  
 KMC = Kanifing Municipal Council  
 UIBC = Unbound iron-binding capacity  
 IL6 = Interleukin 6  
 IL22 = Interleukin 22  
 sTfR = Soluble transferrin receptor  
 CRP = C-reactive protein  
 AGP = Alpha 1-acid glycoprotein  
 HIV = Human immunodeficiency virus  
 TB = Tuberculosis  
 EDTA = Ethylenediaminetetraacetic acid  
 V1 = Venous bleed 1  
 V2 = Venous bleed 2  
 IM = Intramuscular  
 ELISA = Enzyme-linked immunosorbent assay  
 ID = Identification  
 CRF = Case report form  
 eCRF = Electronic case report form  
 ANOVA = Analysis of variance  
 IDE = Integrated development environment  
 CyTOF = Cytometry by time of flight  
 STAT3 = Signal transducer and activator of transcription 3  
 TLR = Toll-like receptor  
 SOP = Standard operating procedure  
 OD = Optical density  
 WBC = White blood cell  
 LPS = Lipopolysaccharide

## Introduction

### Neonatal infections – challenges in low-income settings

Neonatal infection is the third largest cause of death in children under-five worldwide and is an ongoing major global public health challenge (Sustainable Development Goal 3.2)<sup>1</sup>. Between 1990 and 2016 maternal and under-five child mortality has decreased by half<sup>2</sup>. However, the proportion of neonatal

deaths among under-five deaths increased from 37% (1990) to 44% (2013)<sup>3,4</sup>. Today, approximately 2.8 million children die annually during the neonatal period – the first 28 days of life. Of these, 73% die within the first six days of life<sup>3</sup>. An increasing proportion of child deaths are in sub-Saharan Africa<sup>5</sup>, with 60–80% of newborn deaths occurring in low birthweight (LBW) neonates (<2500g at birth)<sup>6</sup>. 95% of all LBW neonates are born in low-income countries<sup>7</sup>. However, the situation is likely to be worse than documented, as neonatal deaths in developing countries are commonly under reported and the records commonly contain errors<sup>8,9</sup>. It is estimated that about one third of deaths in the first month of life, are caused by infections including bacterial sepsis, meningitis, pneumonia, neonatal tetanus, and diarrhoea<sup>10</sup>.

Evidence is lacking on the aetiology of neonatal infections in developing countries, especially from community settings<sup>11,12</sup>. However, the limited data suggests that *Klebsiella* species, *Escherichia coli*, and *Staphylococcus aureus* are common causes of early onset sepsis (EONS)<sup>13–15</sup>. The available antibiotic susceptibility data suggests that pathogens associated with neonatal sepsis in developing countries are often resistant to WHO-recommended empiric antibiotics<sup>13,16</sup>. Antibiotic resistance has emerged with potency over the last few decades due to a multitude of complex reasons. Antibiotic overuse, inappropriate prescribing, inadequate diagnostics, extensive agricultural use, availability of few new antibiotics, and the ease of transportation of resistant bacteria are among the factors contributing to the rise. Equally, bacteria have the ability to rapidly mutate (with or without drug selection pressure) and horizontally transfer genetic material between species (i.e. non-human pathogens) of bacteria<sup>17,18</sup>. Neonates are particularly at risk from antibiotic resistant organisms because they generally succumb before alternative antibiotic regimes can be tried.

Diagnosis of neonatal sepsis with high specificity remains challenging in developing countries. A widely used tool developed by the World Health Organization Young Infants Clinical Signs Studies (YICSS), which includes seven clinical signs to aid diagnosis, has only a 85% sensitivity and 75% specificity for severe bacterial infection during the first week of life<sup>19</sup>. Microbiological identification of a pathogen isolated from blood cultures often has higher specificity, but microbiological laboratory facilities are frequently lacking in low-income settings<sup>19</sup>. With this all in mind, there is an immediate need to improve our understanding of neonatal blood-borne infections and develop novel therapies that could enhance immunological protection possibly via boosting innate immune mechanisms.

### Nutritional immunity

Iron is critical for the human host and most pathogens. Iron is one of the most important factors in the host-pathogen battle for resources. Bacteria and other pathogens have evolved a wide variety of mechanisms to acquire iron from the nutrient rich host (e.g. siderophores and iron specific channels)<sup>20</sup> to aid growth and virulence, with a number of iron acquisition genes concentrated on high pathogenicity islands<sup>21</sup>.

Nutritional immunity describes the normal physiological innate processes used by the host to combat infection by limiting nutrient availability. Key among these processes is the ability to rapidly decrease the circulating concentration of iron (and other transition metals) in response to an infection<sup>22</sup>. The hypoferrremia of inflammation is mediated by the hormone, hepcidin. Research completed in 2000–2001 by three independent research groups led to the discovery of the hepcidin hormone, and the important function it plays in many aspects of iron metabolism<sup>23–25</sup>. Hepcidin is now understood to be the master regulator of iron homeostasis. Unlike any other micronutrient, iron is regulated by a hormone that responds to both infection and nutritional status. The host inflammatory mediators, IL6<sup>26</sup>, IL22<sup>27</sup> and Type 1 interferon<sup>28</sup>, have been found to increase transcription of hepcidin through several Toll-like receptor (TLR) ligands<sup>29</sup> and STAT3 signalling<sup>30,31</sup> resulting in decreased systemic iron concentrations in the circulation. This multifaceted mechanism limits nutrient availability to extracellular invading microorganisms<sup>32</sup>. The system is well documented in mouse models<sup>33–35</sup>, but less so in human studies. However, it is clear that humans with excessive levels of serum iron (e.g. due to hemochromatosis) are predisposed to infection with iron-dependent species of bacteria<sup>36,37</sup>.

### Neonatal hypoferrremia

Although iron metabolism in adults and older children is well studied, the kinetics of iron handling in the early neonatal period, a time of intense physiological change, are poorly understood<sup>38</sup>. Childbirth results in a neonate moving from a semi-allogeneic, protected and nearly sterile environment to one that is abundant in a diverse array of microbes. The delivery process is the initial focal point for the mass bacterial colonisation of the skin and gastrointestinal tract of the neonate<sup>39,40</sup>. Neonates are known to have very low levels of immunological memory and possess an immature immune system<sup>41</sup>. Post-natal iron metabolism in neonates is controlled by an array of different signals, such as hypoxia, erythropoietic drive, maternal and foetal iron stores<sup>42</sup>. A number of studies have investigated serum iron, transferrin saturation (TSAT), ferritin and haemoglobin levels at the time of birth using cord blood as a proxy for early neonatal blood<sup>43,44</sup>. A recent prospective study showed neonates born preterm compared to early-term had higher serum iron concentrations in umbilical blood, which was inversely correlated with levels of serum hepcidin<sup>45</sup>. A similar study has also shown that small-for-gestational-age neonates and neonates born by elective caesarean have lower levels of hepcidin<sup>44</sup>. Previous work has shown that serum iron and TSAT decreases between birth and the first 6–12 hours post-partum in full term, healthy vaginally delivered newborns<sup>46,47</sup>.

The study described here will shed light on the effects of prematurity and birthweight on body iron distribution immediately after birth and during the first week of life. Free ferric and ferrous iron (i.e. transferrin bound iron), haem-based iron molecules and their chaperone proteins (haem-hemopexin and haemoglobin-haptoglobin) will also be investigated.

### Study objectives

The primary study objective is to characterize in detail how full term, preterm and low birthweight neonates modulate serum

iron in the first 24 hours of life. We hypothesize that premature and/or low birthweight babies have a defect in their ability to sequester iron at 6–24 hours after birth in comparison to full term neonates with normal birthweight.

The secondary objectives are:

- I. Characterise how iron metabolism, handling and recycling differs between full term, preterm and low birthweight neonates at birth and during the first 24 hours of life.
- II. Describe iron metabolism, handling and recycling in full term neonates at birth and during the first 7 days of life.
- III. Determine if sera from preterm and low birthweight neonates supports a greater level of *ex-vivo* growth of microorganisms that are common causes of neonatal sepsis in Africa and The Gambia (*Staphylococcus aureus*, *Klebsiella pneumoniae*, *Escherichia coli*, Group B *Streptococcus*, *Streptococcus pneumoniae* and *Salmonella enterica* serovar Typhimurium (*S. Typhimurium* hereafter)) in comparison to sera from full term, normal birthweight neonates.
- IV. Characterize frequencies and functionality of neutrophils, monocytes, dendritic cells, NK cells, B cells and T cells (D8 and CD4) in cord blood from full term, premature and low birthweight neonates.

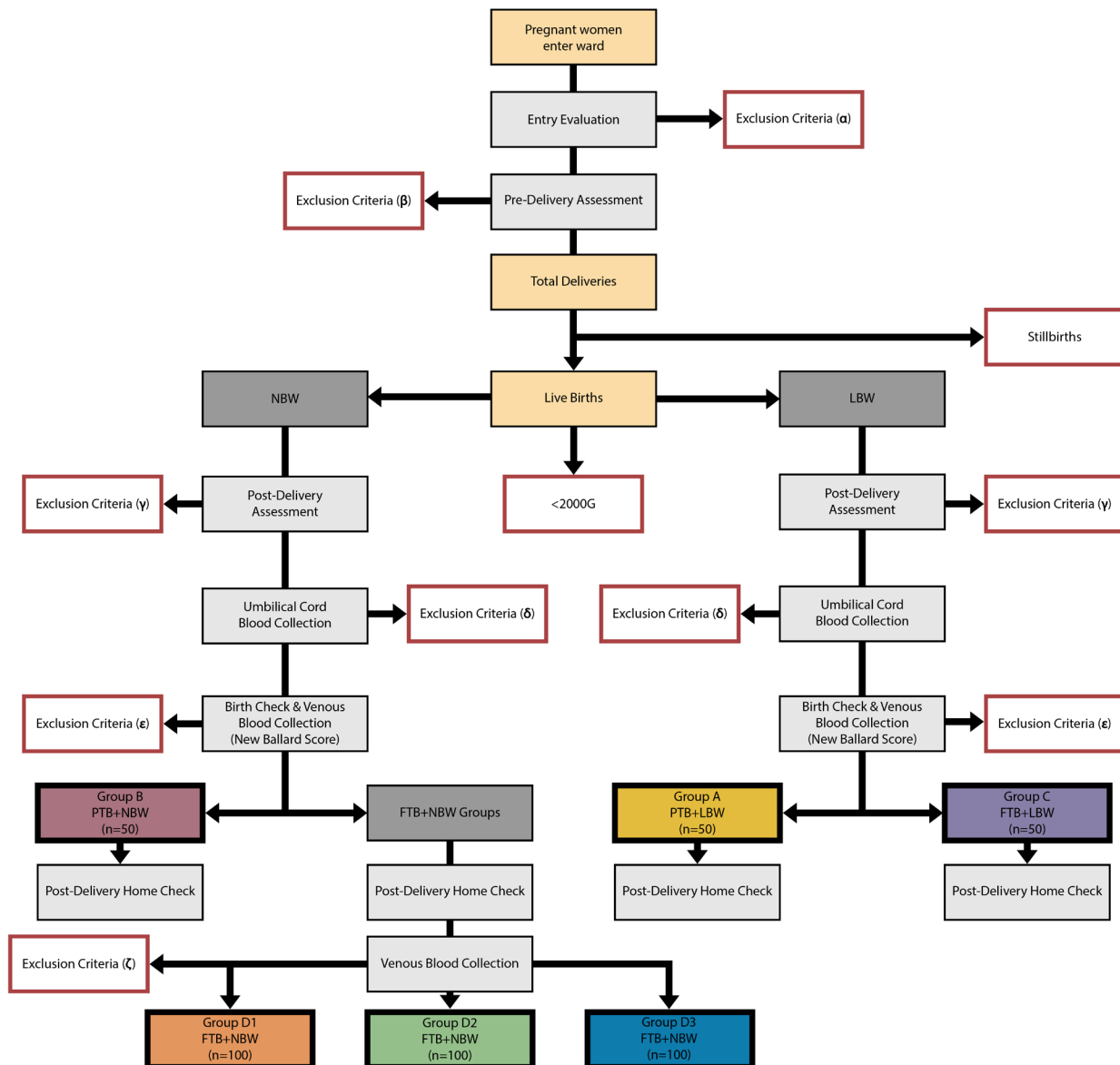
### Protocol

#### Study site

Study participants will be recruited from Kanifing General Hospital (formally Serrekunda General Hospital), in the Kanifing region of The Gambia, West Africa. Serrekunda is a large town, forming a peri-urban area with a population of around 340,000, and is 13km to the southwest of the capital, Banjul. Serrekunda was originally made up of nine villages that have merged into a sprawling urban area. Annually, Kanifing General Hospital (KGH) provides antenatal care to 500–700 pregnant mothers. Mothers receiving antenatal care at other local health-care facilities increase the total number of births at the hospital to 3500–4500 per year. The percentage of these that are live, low birthweight neonates (<2.5kg) is approximately 10%. Specimen samples will be subjected to primary processing on-site at KGH, followed by transport to Medical Research Council Unit The Gambia at LSHTM (MRCG) for storage and analysis.

#### Participants

In total, 450 healthy newly born neonates will be identified during delivery at the Kanifing General Hospital Maternity Ward (Figure 1) starting in July 2017. Pregnant mothers must be over the age of eighteen years. After informed consent is obtained, neonates who meet the inclusion criteria will be enrolled into the study. For inclusion in the study, neonates must be healthy, medically stable, greater than 32 weeks gestational age and weigh more than 2000g. To be considered preterm (PTB) the neonates will be < 37 weeks gestational age (assessed by New Ballard Score<sup>48</sup>) and ≥ 32 weeks gestational age. All neonates with a gestational age ≥37 weeks will be considered full term (FTB). To be considered low birthweight (LBW) the



**Figure 1. Main study flow chart of all study procedures and exclusion criteria.** Group A will contain neonates characterised by preterm birth and low birthweight (PTB+LBW); Group B will contain neonates characterised by preterm birth and normal birthweight (PTB+NBW); Group C will contain neonates characterised by full term birth and low birthweight (FTB+LBW); Groups D1, D2 and D3 will all contain babies characterised by full term birth and normal birthweight (FTB+NBW). In this study, preterm is defined <37 weeks gestation and low birthweight is <2500g. Exclusion criteria (a): Father refused, mother refused, family/escort refused, communication not possible or mother with severe disabilities. Exclusion criteria (b): Antibiotics or antimalarials given before delivery (<24 hours), referred to tertiary level health facility, absconded, known HIV-positive, severe pre-eclampsia, receiving TB treatment, antepartum haemorrhage, recent blood transfusion (within the last month), no foetal heartbeat, mother <18 years, refusal, recruited to another study and emergency caesarean section. Exclusion criteria (y): Recruited to another study on-site, refusal, blood transfusion given in labour, antibiotics or antimalarials given during labour, neonate requires resuscitation (1 min APGAR), neonatal weight <2000g, neonate born breech, neonate born via vacuum delivery, neonate born caesarean section, foetal stillbirth, macerated stillbirth and major congenital malformations. Exclusion criteria (d): Failed cord blood collection (serum tubes), failed cord blood collection (EDTA), cord blood processed >3 hours, neonate requires resuscitation (10 min APGAR), absconded and route 2B refusal. Exclusion criteria (e): Mother birth check refusal, father birth check refusal, family escort birth check refusal, mother <18 years, recruited to another study on-site, antibiotics or antimalarials given to mother before delivery (<24 hours), neonate has had surgery, neonates sibling twin was recruited, neonate given antibiotics (other than tetracycline eye ointment), neonate given iron supplementation, neonatal sickness (tone, activity, feeding, heart rate, respiratory rate, abnormal anterior fontanelle), neonatal temperature (<36.5°C or >37.5°C), major congenital malformations (neonate), New Ballard Score (<32 weeks), failed V1 (serum), failed V1 (EDTA), failed V1 (both EDTA and serum), mother V1 bleed refusal, father V1 bleed refusal, and family/escort V1 bleed refusal. Exclusion criteria (z): neonatal sickness (tone, activity, feeding, heart rate, respiratory rate, abnormal anterior fontanelle), neonatal temperature (<36.5°C or >37.5°C), neonate has had surgery, neonate given antibiotics (other than tetracycline eye ointment), neonate given iron supplementation, failed V2 bleed, Mother community/V2 bleed refusal, father community/V2 bleed refusal, and family community/V2 bleed refusal.



neonates will weigh < 2500g. All neonates weighing  $\geq 2500$  g will be considered normal birthweight (NBW).

The study groups are:

Group A (PTB+LBW): Neonates who are both preterm and low birthweight.

Group B (PTB+NBW): Neonates who are preterm and normal birthweight.

Group C (FTB+LBW): Neonates who are full term but low birthweight.

Group D (FTB+NBW): Neonates who are full term and normal birthweight.

In addition to the main study, 300 FTB neonates of the 450 neonates will also be included into a sub-study, which aims to describe serum iron markers in full term babies (Group D, FTB+NBW only) over the first week of life.

### Study design

This is a proof-of-concept, observational cohort study (Groups A, B, C and D) with an embedded short prospective cohort study (Group D divided into D1, D2 and D3).

### Entry evaluation

**Consent and enrolment.** There are two routes into the study enrolment (Figure 2). Pregnant mothers who are receiving antenatal care on-site at KGH, will be approached at an antenatal visit and voluntarily sensitised to the study requirements and protocol (Route 1). Pregnant women, who are sensitised will not be required at that point to give written or verbal consent. This group will be provided with study information sheets and encouraged to discuss study participation with their family. When the pregnant woman returns to KGH Maternity Ward to deliver (some mothers will choose to deliver at other healthcare facilities), she will be asked to read the full study information sheet (or have it read to her by a study nurse if she is not literate) and provide formal written consent to the study involvement for their neonate (see Extended data<sup>49–51</sup>).

Route 2 will provide an alternative route of enrolment for pregnant mothers, that would like their neonate to be part of the study but have been receiving antenatal care at another facility before delivering at KGH maternity ward. In route 2, healthy, pregnant women will enter the KGH ward to deliver and will be approached to provide written formal consent to umbilical cord blood collection and storage only. No testing or laboratory processes will be conducted on their sample, until full study consent is gained post-delivery. The cord blood sample will be stored at 4°C (within the maternity ward), until the mother's pain and discomfort subside (2–6 hours post-delivery). At this point, the mother and/or father will be invited to provide written formal consent on full study enrolment. If, at this point, mother and/or father refuse full study consent post-delivery, the previously collected personal information and umbilical cord sample will be safely discarded.

### Pre-delivery screening

In both Route 1 and Route 2 enrolment, mothers must provide written consent before assessment of personal information (antenatal card) and questioning can begin. After consenting, mothers will be asked for their demographic information and their personal contact details. Pregnant mothers will be excluded from the study if they are below the age of 18 years, have no foetal heartbeat detected upon admission, known to be HIV-positive, in receipt of *Mycobacterium tuberculosis* therapy, taken antibiotics in the last 24 hours, had a blood transfusion in the last month, suffering from severe pre-eclampsia or antepartum haemorrhage, or in another research study. Mothers can refuse to be part of the study at any stage of the study protocol. Pregnant women that are referred at this point to a tertiary level healthcare facility, will be excluded from the study.

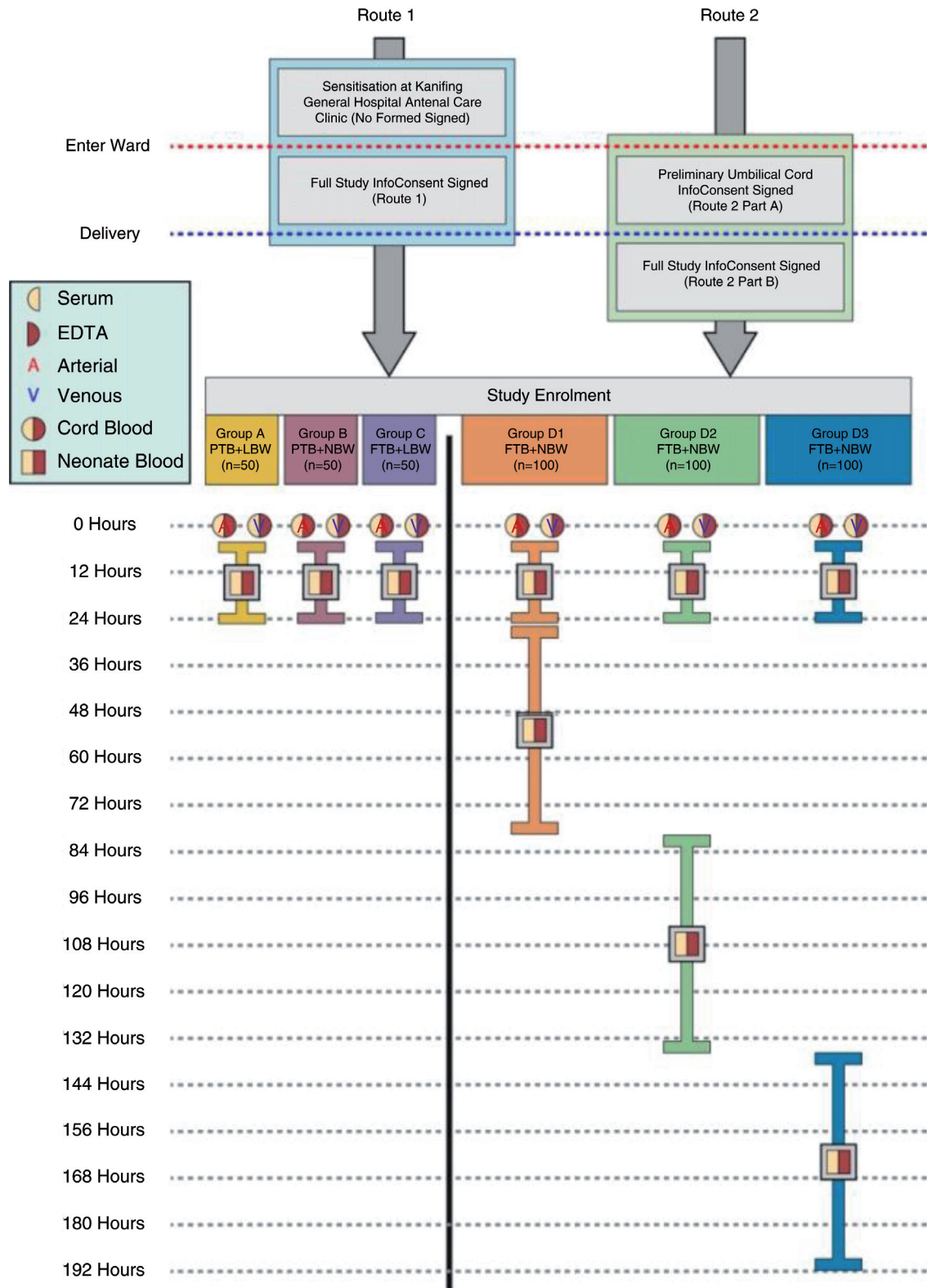
### Delivery procedures, post-delivery screening and umbilical cord blood collection

**Delivery procedures and screening.** Study nurses will assist clinical KGH maternity ward staff in the delivery process and collect data via electronic case report form (eCRF) on their designated study tablets. Neonates will be excluded at the delivery stage of the study for the following reasons: major congenital malformations (not including polydactylism), blood transfusions given to mother or neonate, severe birth asphyxia (requiring resuscitation), neonates born via breech, vacuum or via caesarean section, or a birthweight <2000g. After the delivery stage of the study protocol, neonates can be excluded from the study following the detection of infection or illness (information gained from full blood count analysis or review of systems). Neonates will also be removed from the study protocol, if medication is given (not including intramuscular vitamin K, tetracycline eye ointment or any immunisations). All medication that is given to mothers and neonates will be recorded. Mothers will be able to refuse study participation at any stage. Mothers that deliver multiple newborns will only be invited to consent and enrol one of their neonates into the study.

**Umbilical cord blood collection.** Once the neonate is fully delivered, one-minute delayed cord clamping will be used (following World Health Organisation (WHO) policy<sup>52</sup>). During the one-minute delay, the one-minute APGAR score will be conducted. If the neonate requires resuscitation, the neonate will be excluded from the study. After the umbilical cord has been removed and cleaned, a trained study nurse will identify the umbilical arteries and the umbilical vein. Blood will be collected from both. The tubes will be placed in the cool box for 1–3 hours before transfer to the study laboratory for primary processing. If the mother is enrolled by route 2, the mother will be asked to provide written consent to full study recruitment before the sample is sent for primary laboratory processing.

### Hospital assessment and 1<sup>st</sup> venous blood draw

**Hospital health assessment (study recruitment and group allocation).** At 6–24 hours post-delivery, recruited mothers and their neonates will be invited to a private consultation with the study research clinician. Further demographic data will



**Figure 2. Neolnnate Study enrolment route and blood draw design.** Group A contains neonates characterised by preterm birth and low birthweight (PTB+LBW); Group B contains neonates characterised by preterm birth and normal birthweight (PTB+NBW); Group C contains neonates characterised by full term birth and low birthweight (FTB+LBW); Groups D1, D2 and D3 all contain babies characterised by full term birth and normal birthweight (FTB+NBW).

be collected, along with a photograph of the antenatal card to gather gestational age data (fundal height, last menstrual period and ultrasound), mother's last haemoglobin level before delivery (dated), known sickle cell status, neonate immunisations, and medication given to the mother (pre, during and post-delivery) and the neonate. A complete review of systems of the mother and neonate plus anthropometric data on the newborn will then be collected. Neuromuscular and physical maturation of each neonate will be assessed using the New Ballard Score<sup>48</sup>.

Neonates will be excluded if they score less than 32 weeks of gestation. From this assessment, the neonate will be assigned to a specific study group. If the neonate is allocated to the Group D (FTB+NBW) group, the neonate will be allocated to a randomised bleed group ( $\geq 24$  hours -  $< 80$  hours (Group D1);  $\geq 80$  hours -  $< 136$  hours (Group D2); and  $\geq 136$  -  $< 192$  hours (Group D3). Failure to meet the inclusion criteria at this stage of the study protocol, will result in exclusion from the study.

**1<sup>st</sup> venous blood draw (all neonates).** A blood sample will be collected from all neonates that have passed the inclusion criteria in the hospital health assessment. Immediately after the health assessment, a venous blood draw will be performed (6–24 hours post-delivery). PTB and/or LBW neonates will donate 2ml of venous blood. FTB+NBW neonates will donate 3.5ml of venous blood. All samples will reach the laboratory within three hours post collection for primary processing.

**Community health assessment and 2<sup>nd</sup> venous blood draw**  
**Community health assessment.** Study nurses will visit all mothers or enrolled neonates at their homes at least once. At that visit, a physical examination of the neonate will be completed. The following information will also be collected: neonatal immunisation history, a complete review of systems of the mother and baby, and any medication given to the mother or neonate since delivery. Mothers will also be provided with health education and study contact details (should the neonate become unwell).

**2<sup>nd</sup> venous blood draw Group D (FTB+NBW) only.** At this point, if the mother and neonate are deemed to have passed the screening process and the neonate is in Group D, then the neonate will have its second and last venous blood draw (3.5ml). All samples will reach the laboratory within three hours post collection for primary processing.

### Laboratory evaluations

**Blood samples.** Whole blood samples will be assessed for: full haematology panel (using a Medonic M20M GP), glucose-6-phosphate dehydrogenase deficiency and sickle trait. All serum samples collected will be assessed by ELISA for the following: IL6, IL22, free haem, hepcidin, hemopexin, lipocalin-2, lactoferrin, and foetal haemoglobin. Additionally, serum ferritin, serum iron, UIBC, soluble transferrin receptor (sTfR), transferrin, C-reactive protein (CRP), haptoglobin, and alpha 1-acid glycoprotein (AGP) will be assessed using a fully automated biochemistry analyser (Cobas Integra 400 plus) Additionally, umbilical WBC will be processed and analysed for exploratory secondary analysis 4.

**Bacterial growth assays.** *Ex vivo* growth of bacteria (including clinical and laboratory isolates of *Staphylococcus aureus*, *Klebsiella pneumoniae*, *Escherichia coli*, *Enterobacter* spp., *Enterococcus* spp., and *Salmonella Typhimurium*) in participant serum as in Cross *et al.* (2015)<sup>53</sup> will be performed.

### Study outcomes

The primary outcome variables will be TSAT (transferrin saturation) and serum iron.

The secondary outcome variables will be hepcidin; hemopexin; haptoglobin; IL22; free serum haem and haemoglobin; foetal haemoglobin; lactoferrin; lipocalin-2; IL6; C-reactive protein; alpha-1-acid glycoprotein; transferrin concentration; soluble transferrin receptor; unbound iron-binding capacity; ferritin; haemoglobin; WBCs types and numbers in cord blood samples and *ex vivo* bacterial growth.

### Data entry, handling, storage and security

All protocol-required field data will be captured electronically on an electronic eCRF or a paper case report form (CRF) that will be completed for each included participant. After giving written consent the pregnant women will be given a study identification number, which will be used in all future datasets for subject anonymity. Field data will be collected verbally and from antenatal cards by study nurses. Collected data will be entered in real time using eCRFs developed on top of a REDCap (Research Electronic Data Capture) database and published on Samsung Galaxy Tab 3 SM-T111 handheld devices. Collected data will be transported to the database via a direct secure connection over the 4G mobile network. Laboratory related data will be extracted directly from laboratory equipment and uploaded to the database. Any data collected on the paper format will be double entered by a trained data entry clerk. The local co-investigator will review all forms and identify any errors prior to data entry or to marking data as complete. The study data will also be validated through automated and manual validation methods implemented in the study database application system. The study database will be custom-developed. All paper CRF will be stored in a locked file archive. Electronic data will be stored on the local dedicated server maintained at MRCG. The study will be conducted in compliance with Good Clinical Practice. Study personal security measures will include controlled access limited to authorised users only, physical security, remove identifiable information (anonymization), avoidance of third-party cloud storage and password protection.

### Sample size and power

This study will target recruitment of 150 “exposed” neonates which will include a target of 50 neonates in each Group A, B and C. 300 neonates will be recruited for Group D (“unexposed”). The study will have constraints from time, budget, loss to follow up, haemolysis during sample collection, insufficient blood volume and the distribution of new births in each group at the Kanifing General Hospital.

Based on this, we have run simulations (Stata/IC 15.1) to calculate the power to detect a range of differences comparing groups for example Groups D and A with respect to the primary



outcomes TSAT and serum iron. We did not calculate power for the secondary outcomes, which are considered exploratory. The simulation was run using a linear regression model assuming a lognormal distribution for the response variables TSAT and serum iron levels 6–24 hours after birth. Data from a previous study (Prentice S, personal communication) was used to obtain mean and SD estimates for TSAT and serum iron both at baseline and 6–24 hours after birth. The predictor variables were the Groups (A–D) with Group D as the reference. The model was adjusted for the baseline (cord blood levels). We also examined the power assuming a normal distribution for TSAT (i.e. without log transformation). The significance level considered was 0.05 and the simulation was run for 100000 iterations. This process was repeated for the following four different sample size scenarios which we refer to as *N1*, *N2*, *N3* and *N4* respectively:

- N1*) Group A=Group B=Group C =50 neonates
- N2*) Group A=Group B =25 neonates and Group C =50 neonates
- N3*) Group A=Group B =10 neonates and Group C =50 neonates
- N4*) Group A=Group B =50 neonates and Group C =10 neonates

For all the above four cases, *D*=300.

The simulation results for the baseline adjusted model with log transformation show that for sample size scenario *N1*, the minimum mean differences that can be detected with 80% power were about 4% and 2.5  $\mu\text{mol/L}$  for TSAT and serum iron respectively (Figure 3A and 3B). These correspond to effect sizes of 0.35 and 0.39 respectively. The power drops substantially if smaller numbers were to be recruited as in scenarios *N2* (*A*=*B*=25) and *N3* (*A*=*B*=10). Under *N2* and *N3*, the minimum mean differences that can be detected with 80% would increase to about 5.8% and 9.1% for TSAT (Figure 3A) and 3.3  $\mu\text{mol/L}$  and 5  $\mu\text{mol/L}$  for serum iron (Figure 3B). The results for scenario *N4* can be considered as subset of *N1*–*N3* by rearranging Groups A, B and C.

### Statistical analysis

The primary research objective is to examine if preterm and/or low birthweight neonates (“Exposed”) have a reduced ability to sequester iron at 6–24 hours after birth in comparison to full term neonates with normal birthweight (“Unexposed”)?

We hypothesize that FTB+NBW (Group D) neonates on average will have lower values of TSAT and serum iron compared to “Exposed” (PTB or LBW babies) (Figure 4). Initially, we will analyse all “Exposed” (Groups A+B+C) vs “Unexposed” (Group D). Each neonate will be further classified by his or her gestational age (premature vs. full term) and birthweight (low vs. normal) in a 2x2 table (Table 1). Linear regression models will be used in order to evaluate the difference in mean between each Group A–C and D; that is where *D* will be the reference group. TSAT and serum iron levels will be log transformed before fitting the models (if necessary). Both the unadjusted and adjusted (for the cord blood level) mean differences together with the 95% CI will be calculated.

In the second stage of analysis, we will assess the effect of potential confounding variables using the regression models. Covariates to be considered include the specific time of measurement, demographic and health variables. The time effect may not be linear, and this will be investigated in the further regression models. To reduce the effects of multiple testing, data analysis will be driven by a predefined primary study hypothesis. Any exploratory analyses conducted (in the absence of predefined study hypotheses) will be considered hypothesis-generating, rather than confirmatory. In order to reduce the levels of missing and inaccurately entered data into the database, all clinical, demographic and laboratory data will be entered in real time via electronic data capture, with automated and manual validation methods implemented. The study design does not provide for the recruitment of equal numbers of subjects in each month of the year (or during the dry vs wet seasons). The Gambia has a higher birth rate during the months of September–December in comparison to other months<sup>54</sup>.

In order to remove this potential source of bias, we will adjust for month of birth and/or season in the regression analysis. If the missing data rates is more than 5%, we will consider imputation. The follow-up duration is relatively short. Thus, we expect little bias from loss to follow-up. If loss to follow-up rate is considerably different between groups, we will perform sensitivity analyses to examine the robustness of results. We will also consider sensitivity analysis fitting a multivariate regression model where the main outcomes of interest (including TSAT, serum iron and hepcidin) will be jointly regressed to the same set of predictors.

The analysis for the secondary objectives are described below:

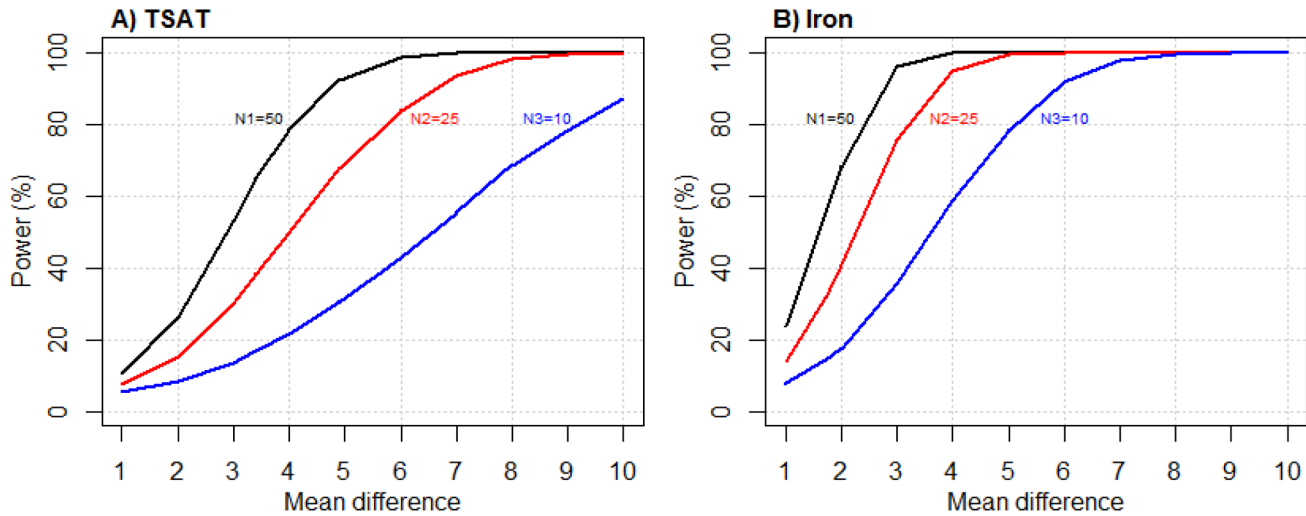
### I. Characterise how iron metabolism, handling and recycling differs between full term, preterm and low birthweight neonates at birth and during the first 24 hours of life.

A similar strategy will be used as for the primary objective. Regression modelling will be used to evaluate the difference in means between each Group A–C and D; D will be the reference (“Unexposed”) group. The effects of potential confounding variables will also be assessed using further regression modelling.

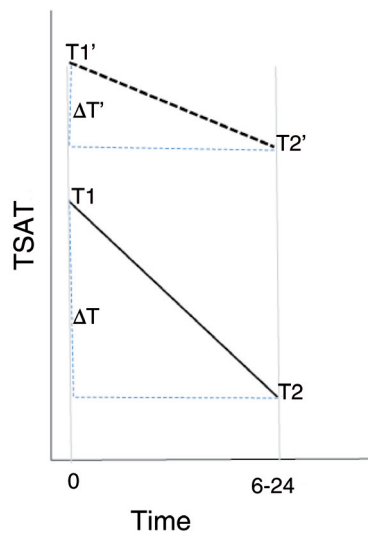
### II. Describe iron metabolism, handling and recycling in full term neonates at birth and during the first 7 days of life.

Analysis of the longitudinal data will involve generalised estimating equations incorporating time of measurement. We will include spline terms at each time point to evaluate the change in the outcomes (all primary and secondary outcome parameters) over time during the transition period from cord to 24–79; 80–135; 136–192 hours after birth. Note that this will only include data collected from Group D neonates and will not be a comparison between Groups A–C and D neonates.

### III. Determine if sera from preterm and low birthweight neonates supports a greater level of *ex-vivo* growth of



**Figure 3.** Estimated power to detect a given difference between Groups A vs D based on simulation using a linear regression model adjusted for baseline for three sample size scenarios. N1 (Group A=Group B=Group C=50 neonates); N2 (Group A=Group B=25 neonates and Group C=50 neonates); N3 (Group A=Group B=10 neonates and Group C=50 neonates); N4 (Group A=Group B=50 neonates and Group C=10 neonates).



**Figure 4.** An example of hypothetical scenario for TSAT values between the groups to be compared. In this example: (i) Time 0 refers to average cord blood levels (ii) Time 6-24 refers to the mean level in the 6-24 hour period after birth. (iii) T1, T2, represent TSAT in 1 and 2 above and  $\Delta T = T2 - T1$  for full term, normal birthweight (Group D) (iv). T1', T2' and  $\Delta T' = T2' - T1'$  same as above but for the premature, low birthweight (Group A). Hypothesis:  $H_0$ :  $T2 = T2'$  vs.  $H_A$ :  $T2 \neq T2'$ .

microorganisms that are common causes of neonatal sepsis in Africa and The Gambia (*Staphylococcus aureus*, *Klebsiella pneumoniae*, *Escherichia coli*, *Enterobacter* spp., *Enterococcus* spp., and *S. Typhimurium*) in comparison to sera from FTB+NBW neonates.

The bacterial growth will be analysed in a similar method as described in Cross *et al.* (2015)<sup>53</sup> in order to determine if

**Table 1.** Four combinations in total for exposure: a: Pre, Low; b: Pre, Normal; c: Full, Low and d: Full, Normal.

		Birthweight (BW)	
		Low	Normal
Term	Pre	A	B
	Full	C	D

changes in iron availability modulate the growth. Growth assays will be fitted to a standard form of the logistic equation:

$$N_t = \frac{K}{1 + \left(\frac{K - N_0}{N_0}\right)e^{-rt}}$$

Here, the population size at the beginning of the growth curve is given by  $N_0$ . The carrying capacity is given by  $K$ . The intrinsic growth rate of the population is  $r$ . We will generate the best fitting values of  $K$ ,  $r$  and  $N_0$  for the growth curve data. Additionally, for each bacterium, we compare the time at which the population density reaches  $\frac{1}{2}K$  (inflection point), the fastest possible generation time (doubling time) and the area under the logistic curve obtained by taking the integral of the logistic equation. This will be used to assess growth curves from different sample types (Cord vs V1) and between the four study groups.

**IV. Characterize frequencies and functionality of neutrophils, monocytes, dendritic cells, NK cells, B cells, T cells (D8 and CD4) in cord blood from full term, premature and low birthweight neonates.**

Exploratory analysis will be conducted using linear regressions modelling.

Statistical analyses will be performed using **STATA** (StataCorp. 2017. Stata Statistical Software: Release 15. College Station, TX: StataCorp LLC); **R** (R Foundation for Statistical Computing, Vienna, Austria.) and **Data Desk** (Data Description Inc Ithaca NY). All files used will have an accompanying data dictionary. Annotated STATA do-files or R files will be used to describe any data transformations and statistical tests used.

### Dissemination of findings

The study results will be published in relevant peer-reviewed journals and key findings will be presented at international scientific meetings. Data sharing will be in agreement with the sponsor policy on research data sharing and with the Bill & Melinda Gates Foundation Global Access requirements.

### Study status

The study is in the data collection phase.

### Discussion

Humans and bacteria are involved in an on-going tug of war over iron. Each side has evolved complicated and varied iron-acquisition mechanisms in an effort to turn the tide of war in their own favour<sup>55</sup>. Nutritional immunity describes the processes by which the human host tries to starve invading organisms of nutrients, especially iron.

This study aims to determine if premature and low birthweight babies have a defect in their ability to sequester iron during the first 24 hours of life. The study design will produce a detailed and extensive picture of iron metabolism in neonates. To our knowledge, no other study has tried to analyse such a large and diverse collection of iron and infection variables in neonates born in Sub Saharan Africa. The study will enrol subjects who are all at an increased risk of neonatal infection, and subsequent sepsis and death.

A potential limitation of this study is the inadequacy of using the New Ballard Score as the only method of gestational aging. Original and New Ballard Score are reported to overestimated gestational age compared to ultrasound and in particular, misclassify preterm infants as term newborns<sup>56</sup>. Additionally, newborn clinical assessments as a whole, tend to underestimate gestational age in growth-restricted neonates<sup>56</sup>. The gold standard of gestational aging is an ultrasound in the first trimester<sup>57</sup>. However, this procedure is rarely correctly completed in this study population. If it is documented on the mother's antenatal records, care will be taken to record it. Limits of the study also include that HIV status, TB status and iron supplementation given are all gained from the antenatal records of the mother. Furthermore, antenatal records will not contain all information on medication given in every mothers' pregnancy. As a result,

care will be made to extensively question participants mother's during verbal one-to-one consultation with our study research clinician.

In conclusion, our overarching study goal is to evaluate the likelihood that novel products designed to induce hypoferraemia (potentially via mini-hepcidins<sup>58</sup>) may be useful in the future for the prevention of neonatal sepsis in high risk babies. This could be produced by a transient redistribution of iron away from the circulation, thus applying a bacteriostatic brake on any bacteria that have crossed into the baby's systemic circulation and hence boosting host survival in vulnerable newborns. We hope this may ultimately help reduce the use of antibiotics in maternal and neonatal wards worldwide.

### Ethical approval

This study has been approved by The Gambia Government/MRC Joint Ethics Committee (no. SCC1525) and Ethics Committee of London School of Hygiene and Tropical Medicine (ref no. 14316). The study procedures will be explained to the neonate's mother/guardians orally or in writing. A neonate is only recruited into the study after the consent form has been signed/thumb printed by the mother/guardian.

This study was registered with clinicaltrials.gov ([NCT03353051](https://clinicaltrials.gov/ct2/show/study?term=NCT03353051)) on 27 November 2017.

### Data availability

#### Underlying data

No data are associated with this article.

#### Extended data

Figshare: Cross *et al.* GatesOpen Research SCC1525v2\_\_NeoInnate\_Participant Info&Consent form Route 1. <https://doi.org/10.6084/m9.figshare.8069195.v4><sup>49</sup>

This project contains the following extended data:

- SCC1525v2\_\_NeoInnate\_Consent form Route 1\_v3 Approved8Nov17.docx (Route 1 consent and information sheet)

Figshare: Cross *et al.* GatesOpenResearch SCC1525v2\_\_NeoInnate\_Consent form Route 2\_Part 1\_ (Umbilical Cord Blood Collection) - Labour Ward\_v1.1-Approved 8Nov17. <https://doi.org/10.6084/m9.figshare.8069246.v1><sup>50</sup>

This project contains the following extended data:

- SCC1525v2\_\_NeoInnate\_Consent form Route 2\_Part 1\_ (UCB Collection) - Labour Ward\_v1.1-Approved 8Nov17.docx (Route 2 consent and information sheet part 1)

Figshare: Cross *et al.* Gates Open Research SCC1525v2\_\_NeoInnate\_Consent form Route 2\_Part 2\_(Post-Delivery) - ANC

Outside SGH v1-Approved 8Nov17. <https://doi.org/10.6084/m9.figshare.8069243.v1><sup>51</sup>

This project contains the following extended data:

- SCC1525v2\_\_NeoInnate\_Consent form Route 2\_Part 2\_(Post-Delivery) - ANC Outside SGH v1-Approved 8Nov17.docx (Route 2 consent and information sheet part 2)

Data are available under the terms of the [Creative Commons Attribution 4.0 International license](#) (CC-BY 4.0).

## Grant information

This study was supported by the Bill & Melinda Gates Foundation [OPP1152353].

*The funding agency had no role in the design and conduct of the study, and will not have any in the collection, management, analyses or interpretation of the data nor in the preparation, review, or approval of the manuscript.*

## Acknowledgements

We thank Kanifing Municipal Council (KMC) and Kanifing General Hospital (KGH) for their support during the study.

## References

- WHO: **WHO | Every Newborn: an action plan to end preventable deaths**. WHO (World Health Organization), 2014. [Reference Source](#)
- United Nations Children's Fund, World Health Organization, W. B. and U. N. **Levels & Trends in Child Mortality. Report 2017. Estimates developed by the UN Inter-agency Group for Child Mortality Estimation**. New York (NY): United Nations Children's Fund. Levels & Trends in Child Mortality. Report 2017. 2017. [Reference Source](#)
- WHO; UNICEF: **UNICEF - Levels & Trends in Child Mortality. Rep.2014**, 2014. [Reference Source](#)
- Lawn JE, Kinney MV, Black RE, *et al.*: **Newborn survival: a multi-country analysis of a decade of change**. *Health Policy Plan*. 2012; **27** Suppl 3: iii6–iii28. [PubMed Abstract](#) | [Publisher Full Text](#)
- United Nations Inter-agency Group for Child Mortality Estimation (UN IGME): **United Nations Inter-agency Group for Child Mortality Estimation (UN IGME), 'Levels & Trends in Child Mortality: Report 2018, Estimates developed by the United Nations Inter-agency Group for Child Mortality Estimation'**. United Nations Children's Fund, New. [Reference Source](#)
- World Health Organization: **Care of the preterm and/or low-birth-weight newborn**. [Updated 2017]. Accessed Dec 2018. [Reference Source](#)
- UNICEF & WHO: **United Nations Children's Fund and World Health Organization, Low Birthweight: Country, regional and global estimates**. UNICEF, New York. 2004; 27. [Reference Source](#)
- Oza S, Cousens SN, Lawn JE: **Estimation of daily risk of neonatal death, including the day of birth, in 186 countries in 2013: a vital-registration and modelling-based study**. *Lancet Glob Health*. 2014; **2**(11): e635–e644. [PubMed Abstract](#) | [Publisher Full Text](#)
- Hill K, Choi Y: **Neonatal mortality in the developing world**. *Demogr Res*. 2006; **14**: 429–452. [PubMed Abstract](#) | [Publisher Full Text](#)
- Lawn JE, Cousens S, Zupan J, *et al.*: **4 million neonatal deaths: when? Where? Why?** *Lancet*. 2005; **365**(9462): 891–900. [PubMed Abstract](#) | [Publisher Full Text](#)
- Huynh BT, Padgett M, Garin B, *et al.*: **Burden of bacterial resistance among neonatal infections in low income countries: how convincing is the epidemiological evidence?** *BMC Infect Dis*. 2015; **15**: 127. [PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)
- Saha SK, Schrag SJ, El Arifeen S, *et al.*: **Causes and incidence of community-acquired serious infections among young children in south Asia (ANISA): an observational cohort study**. *Lancet*. 2018; **392**(10142): 145–159. [PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)
- Waters D, Jawad I, Ahmad A, *et al.*: **Aetiology of community-acquired neonatal sepsis in low and middle income countries**. *J Glob Health*. 2011; **1**(2): 154–70. [PubMed Abstract](#) | [Free Full Text](#)
- Simonsen KA, Anderson-Berry AL, Delair SF, *et al.*: **Early-onset neonatal sepsis**. *Clin Microbiol Rev*. 2014; **27**(1): 21–47. [PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)
- Okomo U: **Neonatal Infections; a hospital-based study in The Gambia examining aetiology and associated maternal Colonisation**. London School of Hygiene & Tropical Medicine. 2018. [Publisher Full Text](#)
- Lubell Y, Ashley EA, Turner C, *et al.*: **Susceptibility of community-acquired pathogens to antibiotics in Africa and Asia in neonates—an alarmingly short review**. *Trop Med Int Health*. 2011; **16**(2): 145–151. [PubMed Abstract](#) | [Publisher Full Text](#)
- Laxminarayan R, Duse A, Wattal C, *et al.*: **Antibiotic resistance—the need for global solutions**. *Lancet Infect Dis*. 2013; **13**(12): 1057–1098. [PubMed Abstract](#) | [Publisher Full Text](#)
- Davies J, Davies D: **Origins and evolution of antibiotic resistance**. *Microbiol Mol Biol Rev*. 2010; **74**(3): 417–33. [PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)
- Young Infants Clinical Signs Study Group: **Clinical signs that predict severe illness in children under age 2 months: a multicentre study**. *Lancet*. 2008; **371**(9607): 135–142. [PubMed Abstract](#) | [Publisher Full Text](#)
- Caza M, Kronstad JW: **Shared and distinct mechanisms of iron acquisition by bacterial and fungal pathogens of humans**. *Front Cell Infect Microbiol*. 2013; **3**: 80. [PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)
- Koczura R, Kaznowski A: **Occurrence of the *Yersinia* high-pathogenicity island and iron uptake systems in clinical isolates of *Klebsiella pneumoniae***. *Microb Pathog*. 2003; **35**(5): 197–202. [PubMed Abstract](#) | [Publisher Full Text](#)
- Drakesmith H, Prentice AM: **Hepcidin and the iron-infection axis**. *Science*. 2012; **338**(6108): 768–772. [PubMed Abstract](#) | [Publisher Full Text](#)
- Krause A, Neitz S, Mägert HJ, *et al.*: **LEAP-1, a novel highly disulfide-bonded human peptide, exhibits antimicrobial activity**. *FEBS Lett*. 2000; **480**(2–3): 147–50. [PubMed Abstract](#) | [Publisher Full Text](#)
- Pigeon C, Ilyin G, Courselaud B, *et al.*: **A new mouse liver-specific gene, encoding a protein homologous to human antimicrobial peptide hepcidin, is overexpressed during iron overload**. *J Biol Chem*. 2001; **276**(11): 7811–7819. [PubMed Abstract](#) | [Publisher Full Text](#)
- Park CH, Valore EV, Waring AJ, *et al.*: **Hepcidin, a urinary antimicrobial peptide synthesized in the liver**. *J Biol Chem*. 2001; **276**(11): 7806–7810. [PubMed Abstract](#) | [Publisher Full Text](#)
- Nemeth E, Rivera S, Gabayan V, *et al.*: **IL-6 mediates hypoferrremia of inflammation by inducing the synthesis of the iron regulatory hormone hepcidin**. *J Clin Invest*. 2004; **113**(9): 1271–1276. [PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)
- Armitage AE, Eddowes LA, Gileadi U, *et al.*: **Hepcidin regulation by innate immune and infectious stimuli**. *Blood*. 2011; **118**(15): 4129–39. [PubMed Abstract](#) | [Publisher Full Text](#)
- Ryan JD, Altamura S, Devitt E, *et al.*: **Pegylated interferon- $\alpha$  induced hypoferrremia is associated with the immediate response to treatment in hepatitis C**. *Hepatology*. 2012; **56**(2): 492–500. [PubMed Abstract](#) | [Publisher Full Text](#)
- Peyssonnaud C, Zinkernagel AS, Datta V, *et al.*: **TLR4-dependent hepcidin expression by myeloid cells in response to bacterial pathogens**. *Blood*. 2006; **107**(9): 3727–32. [PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)
- Wrighting DM, Andrews NC: **Interleukin-6 induces hepcidin expression through STAT3**. *Blood*. 2006; **108**(9): 3204–3209. [PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)
- Verga Falzacappa MV, Vujic Spasic M, Kessler R, *et al.*: **STAT3 mediates hepatic hepcidin expression and its inflammatory stimulation**. *Blood*. 2007; **109**(1): 353–358. [PubMed Abstract](#) | [Publisher Full Text](#)
- Rodríguez R, Jung CL, Gabayan V, *et al.*: **Hepcidin induction by pathogens and**



- pathogen-derived molecules is strongly dependent on interleukin-6. *Infect Immun*. 2014; **82**(2): 745–52.  
[PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)
33. Arezes J, Jung G, Gabayan V, *et al.*: **Hepcidin-induced hypoferremia is a critical host defense mechanism against the siderophilic bacterium *Vibrio vulnificus*.** *Cell Host Microbe*. 2015; **17**(1): 47–57.  
[PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)
34. Michels KR, Zhang Z, Bettina AM, *et al.*: **Hepcidin-mediated iron sequestration protects against bacterial dissemination during pneumonia.** *JCI Insight*. 2017; **2**(6): e92002.  
[PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)
35. Stefanova D, Raychev A, Arezes J, *et al.*: **Endogenous hepcidin and its agonist mediate resistance to selected infections by clearing non-transferrin-bound iron.** *Blood*. 2017; **130**(3): 245–257.  
[PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)
36. Frank KM, Schneewind O, Shieh WJ: **Investigation of a researcher's death due to septicemic plague.** *N Engl J Med*. 2011; **364**(26): 2563–2564.  
[PubMed Abstract](#) | [Publisher Full Text](#)
37. Khan FA, Fisher MA, Khakoo RA: **Association of hemochromatosis with infectious diseases: expanding spectrum.** *Int J Infect Dis*. 2007; **11**(6): 482–487.  
[PubMed Abstract](#) | [Publisher Full Text](#)
38. Collard KJ: **Iron homeostasis in the neonate.** *Pediatrics*. 2009; **123**(4): 1208–16.  
[PubMed Abstract](#) | [Publisher Full Text](#)
39. Dominguez-Bello MG, Costello EK, Contreras M, *et al.*: **Delivery mode shapes the acquisition and structure of the initial microbiota across multiple body habitats in newborns.** *Proc Natl Acad Sci U S A*. 2010; **107**(26): 11971–11975.  
[PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)
40. Houghteling PD, Walker WA: **Why is initial bacterial colonization of the intestine important to infants' and children's health?** *J Pediatr Gastroenterol Nutr*. 2015; **60**(3): 294–307.  
[PubMed Abstract](#) | [Free Full Text](#)
41. Basha S, Surendran N, Pichichero M: **Immune responses in neonates.** *Expert Rev Clin Immunol*. 2014; **10**(9): 1171–84.  
[PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)
42. Lipiński P, Styś A, Starzyński RR: **Molecular insights into the regulation of iron metabolism during the prenatal and early postnatal periods.** *Cell Mol Life Sci*. 2013; **70**(1): 23–38.  
[PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)
43. Lorenz L, Peter A, Poets CF, *et al.*: **A review of cord blood concentrations of iron status parameters to define reference ranges for preterm infants.** *Neonatology*. 2013; **104**(3): 194–202.  
[PubMed Abstract](#) | [Publisher Full Text](#)
44. Lorenz L, Herbst J, Engel C, *et al.*: **Gestational age-specific reference ranges of hepcidin in cord blood.** *Neonatology*. 2014; **106**(2): 133–139.  
[PubMed Abstract](#) | [Publisher Full Text](#)
45. Ru Y, Pressman EK, Guillet R, *et al.*: **Umbilical Cord Hepcidin Concentrations Are Positively Associated with the Variance in Iron Status among Multiple Birth Neonates.** *J Nutr*. 2018; **148**(11): 1716–1722.  
[PubMed Abstract](#) | [Publisher Full Text](#)
46. Szabó M, Vársárhelyi B, Balla G, *et al.*: **Acute postnatal increase of extracellular antioxidant defence of neonates: the role of iron metabolism.** *Acta Paediatr*. 2001; **90**(10): 1167–1170.  
[PubMed Abstract](#) | [Publisher Full Text](#)
47. Sturgeon P: **Studies of iron requirements in infants and children. I. Normal values for serum iron, copper and free erythrocyte protoporphyrin.** *Pediatrics*. 1954; **13**(2): 107–25.  
[PubMed Abstract](#)
48. Ballard JL, Khoury JC, Wedig K, *et al.*: **New Ballard Score, expanded to include extremely premature infants.** *J Pediatr*. 1991; **119**(3): 417–23.  
[PubMed Abstract](#) | [Publisher Full Text](#)
49. Cross J, Jarjou O, Mohammed NI: **Cross *et al.* GatesOpen Research SCC1525v2\_\_Neolnnate\_Participant Info&Consent form Route 1. figshare.** Figure. 2019.  
<http://www.doi.org/10.6084/m9.figshare.8069195.v4>
50. Cross J, Jarjou O, Mohammed NI, *et al.*: **Cross *et al.* GatesOpenResearch SCC1525v2\_\_Neolnnate\_Participant Info&Consent form Route 2\_Part 1\_(Umbilical Cord Blood Collection) - Labour Ward\_v1.1-Approved 8Nov17. figshare.** Figure. 2019.  
<http://www.doi.org/10.6084/m9.figshare.8069246.v1>
51. Cerami C, Cross J, Jarjou O, *et al.*: **Cross *et al.* Gates Open Research SCC1525v2\_\_Neolnnate\_Participant Info&Consent form Route 2\_Part 2\_(Post-Delivery) - ANC Outside SGH\_v1-Approved 8Nov17. figshare.** Figure. 2019.  
<http://www.doi.org/10.6084/m9.figshare.8069243.v1>
52. **WHO | Delayed umbilical cord clamping for improved maternal and infant health and nutrition outcomes.** WHO, 2018.  
[Reference Source](#)
53. Cross JH, Bradbury RS, Fulford AJ, *et al.*: **Oral iron acutely elevates bacterial growth in human serum.** *Sci Rep*. 2015; **5**: 16670.  
[PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)
54. Moore SE, Fulford AJ, Streatfield PK, *et al.*: **Comparative analysis of patterns of survival by season of birth in rural Bangladeshi and Gambian populations.** *Int J Epidemiol*. 2004; **33**(1): 137–143.  
[PubMed Abstract](#) | [Publisher Full Text](#)
55. Skaar EP: **The battle for iron between bacterial pathogens and their vertebrate hosts.** *PLoS Pathog*. 2010; **6**(8): e1000949.  
[PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)
56. Lee AC, Panchal P, Folger L, *et al.*: **Diagnostic Accuracy of Neonatal Assessment for Gestational Age Determination: A Systematic Review.** *Pediatrics*. 2017; **140**(6): pii: e20171423.  
[PubMed Abstract](#) | [Publisher Full Text](#)
57. Benson CB, Doubilet PM: **Sonographic prediction of gestational age: accuracy of second- and third-trimester fetal measurements.** *AJR Am J Roentgenol*. 1991; **157**(6): 1275–1277.  
[PubMed Abstract](#) | [Publisher Full Text](#)
58. Sebastiani G, Wilkinson N, Pantopoulos K: **Pharmacological Targeting of the Hepcidin/Ferroportin Axis.** *Front Pharmacol*. 2016; **7**: 160.  
[PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)

## Chapter 5 – Early postnatal hypoferremia in low birthweight and preterm babies: A prospective cohort study in hospital-delivered Gambian neonates (Main Paper)

---

### Summary of Chapter

**BACKGROUND:** Neonates, particularly those born preterm (PTB) and with low birthweight (LBW), are especially susceptible to bacterial and fungal infections that cause an estimated 225,000 deaths annually globally. Iron is a vital nutrient for the most common organisms causing septicaemia. Full-term babies elicit an immediate postnatal hypoferremia assumed to have evolved as an innate defence. We tested whether PTB and LBW babies are capable of the same response.

**METHODS:** We conducted an observational study of 152 babies who were either PTB (born  $\geq 32$  to  $< 37$  weeks gestational age) and/or LBW ( $< 2500$ g) (PTB/LBW) and 278 term, normal-weight babies (FTB/NBW). Blood was sampled from the umbilical cord vein and artery, and matched venous blood samples were taken from all neonates between 6-24 hrs after delivery. We measured haematological, iron and proinflammatory biomarkers.

**FINDINGS:** In both PTB/LBW and FTB/NBW babies, serum iron decreased 3-fold within 12hrs of delivery compared to umbilical blood ( $7.5 \pm 4.5$  vs  $23.3 \pm 7.1$  ng/ml,  $P < 0.001$ ,  $n = 425$ ). Transferrin saturation showed a similar decline with a consequent increase in unsaturated iron-binding capacity. C-reactive protein levels increased over 10-fold ( $P < 0.001$ ) and hepcidin levels doubled ( $P < 0.001$ ). There was no difference in any of these responses between PTB/LBW and FTB/NBW babies.

**INTERPRETATION:** Premature or low birthweight babies are able to mount a very rapid hypoferremia that is indistinguishable from that in normal term babies. The data suggest that this is a hepcidin-mediated response triggered by acute non-infective inflammation at birth, and likely to have evolved as an innate immune response against bacterial and fungal septicaemia.

## RESEARCH PAPER COVER SHEET - MAIN PAPER

---

Please note that a cover sheet must be completed for each research paper included within a thesis.

### **SECTION A – Student Details**

<b>Student ID Number</b>	LSH158152	<b>Title</b>	Mr.
<b>First Name(s)</b>	James Henry		
<b>Surname/Family Name</b>	Cross		
<b>Thesis Title</b>	Iron and Infection: Neonatal Iron Transition		
<b>Primary Supervisor</b>	Dr Carla Cerami		

If the Research Paper has previously been published please complete Section B, if not please move to Section C.

### **SECTION B – Paper already published**

Where was the work published?

When was the work published?

If the work was published prior to registration for your research degree, give a brief rationale for its inclusion

Have you retained the copyright for the work?*	Choose an item.	Was the work subject to academic peer review?	Choose an item.
--	-----------------	---	-----------------

\*If yes, please attach evidence of retention. If no, or if the work is being included in its published format, please attach evidence of permission from the copyright holder (publisher or other author) to include this work.

### **SECTION C – Prepared for publication, but not yet published**

Where is the work intended to be published?	EbioMedicine
Please list the paper's authors in the intended authorship order:	James H. Cross, Ousman Jarjou, Nuredin Ibrahim Mohammed, Santiago Rayment Gomez, Bubacarr J.B Touray, Andrew M. Prentice, Carla Cerami.
Stage of publication	In press

### **SECTION D – Multi-authored work**

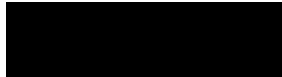


For multi-authored work, give full details of your role in the research included in the paper and in the preparation of the paper. (Attach a further sheet if necessary)

James H. Cross is PhD student with MRCG at LSHTM. He contributed to the development of the protocols, study management, field data collection, development of the data analysis plan, conducting the data analysis and drafting of the manuscript.

## **SECTION E**

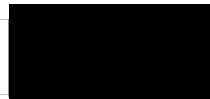
**Student Signature**



**Date**

08/11/2019

**Supervisor Signature**



**Date**

08/11/2019

**Title**

**Early postnatal hypoferrremia in low birthweight and preterm babies: A prospective cohort study in hospital-delivered Gambian neonates**

**Authors**

James H. Cross, BSc<sup>1</sup>, Ousman Jarjou, MBChB<sup>1</sup>, Nuredin Ibrahim Mohammed, PhD<sup>1</sup>, Santiago Rayment Gomez<sup>2</sup>, Bubacarr J.B Touray, BSc<sup>1</sup>, Andrew M. Prentice, PhD<sup>1</sup>, Carla Cerami, MD<sup>1\*</sup>.

**Affiliations**

<sup>1</sup>Medical Research Council Unit The Gambia at the London School of Hygiene & Tropical Medicine, Atlantic Boulevard, Fajara, P.O. Box 273, Banjul, The Gambia

<sup>2</sup>Cardiff University, Cardiff, Wales, CF10 3AT, United Kingdom

**Corresponding Author**

Dr Carla Cerami, email: [ccerami@mrc.gm](mailto:ccerami@mrc.gm)

Medical Research Council Unit The Gambia at the London School of Hygiene & Tropical Medicine, Atlantic Boulevard, Fajara, P.O. Box 273, Banjul, The Gambia

+220 787 5756

**Running Title:**

Neonatal hypoferrremia in premature and low birthweight babies

**Research in context:****Evidence before this study:**

We searched PubMed and Ovid Medline for publications that detail hepcidin, serum iron and TSAT concentrations in umbilical cord blood or venous blood taken in the neonatal period that were published before July 1, 2019 with restriction to publications in the English language. We used the search terms “neonate”, “hepcidin”, “TSAT” and “serum iron”. We excluded studies on non-humans, in medically unstable neonates or studies that did not provide details of gestational age for the study population. We identified 20 studies on hepcidin, 23 on TSAT and 51 on serum iron. Collectively, 59 publications contained research on full-term (FTB) neonates, and 16 on premature (PTB) neonates. Many of the retrieved publications focused on maternal-fetal iron endowment, where umbilical cord blood was used as a proxy for early neonatal blood. Very few publications have documented iron and inflammatory variables in matched umbilical and neonatal blood samples taken in the immediate hours (0-24h) after delivery. Analysis of the retrieved publications leads to the conclusion that neonatal hypoferremia may exist in all neonates, however the evidence is sparse and unclear, particularly in PTB newborns.

**Added value of this study:**

To our knowledge, this observational non-blinded study is the first to assess the independent effects of gestational age and birthweight on iron and infection parameters in healthy Gambian newborns in the first hours of life. Our findings confirm that a very rapid hypoferremia occurs in the early hours of postnatal life with evidence that it is mediated by an increase in hepcidin. Premature and low birthweight babies exhibited almost identical postnatal hypoferremic responses to full-term, normal-weight newborns.

**Implications of all the available evidence:**

There is now clear evidence that neonates, including those with PTB or LBW, elicit a rapid and transient hypoferrremia probably induced, at least in part, by hepcidin. The strength and consistency of this effect indicates that it may have evolved as an innate immune response designed to protect neonates from bacterial septicaemia. This suggests the possibility that a further enhancement or prolongation of hypoferrremia (for instance by mini-hepcidins), might offer an additional tool in the armoury against antibiotic-resistant infections in newborns.

## 5.1 ABSTRACT

**BACKGROUND:** Neonates, particularly those born preterm (PTB) and with low birthweight (LBW), are especially susceptible to bacterial and fungal infections that cause an estimated 225,000 deaths annually globally. Iron is a vital nutrient for the most common organisms causing septicaemia. Full-term babies elicit an immediate postnatal hypoferraemia assumed to have evolved as an innate defence. We tested whether PTB and LBW babies are capable of the same response.

**METHODS:** We conducted an observational study of 152 babies who were either PTB (born  $\geq 32$  to  $< 37$  weeks gestational age) and/or LBW ( $< 2500$ g) (PTB/LBW) and 278 term, normal-weight babies (FTB/NBW). Blood was sampled from the umbilical cord vein and artery, and matched venous blood samples were taken from all neonates between 6-24 hrs after delivery. We measured haematological, iron and proinflammatory biomarkers.

**FINDINGS:** In both PTB/LBW and FTB/NBW babies, serum iron decreased 3-fold within 12hrs of delivery compared to umbilical blood ( $7.5 \pm 4.5$  vs  $23.3 \pm 7.1$  ng/ml,  $P < 0.001$ ,  $n = 425$ ). Transferrin saturation showed a similar decline with a consequent increase in unsaturated iron-binding capacity. C-reactive protein levels increased over 10-fold ( $P < 0.001$ ) and hepcidin levels doubled ( $P < 0.001$ ). There was no difference in any of these responses between PTB/LBW and FTB/NBW babies.

**INTERPRETATION:** Premature or low birthweight babies are able to mount a very rapid hypoferraemia that is indistinguishable from that in normal term babies. The data suggest that this is a hepcidin-mediated response triggered by acute non-infective inflammation at birth, and likely to have evolved as an innate immune response against bacterial and fungal septicaemia.

**Trial registration:** [clinicaltrials.gov](https://clinicaltrials.gov) (NCT03353051). Registration date: November 27, 2017.

**FUNDING:** Bill & Melinda Gates Foundation (OPP1152353).

**WORD COUNT: 247/250**

**KEYWORDS:** Nutritional immunity, iron, transferrin saturation, CRP, hepcidin, neonates, low birthweight, prematurity, sub-Saharan Africa, septicaemia.

## 5.2 INTRODUCTION

Neonatal sepsis is the third highest cause of death globally, accounting for 225,000 deaths each year.<sup>1</sup> It is projected that these numbers will increase as a consequence of the increasing global prevalence of antimicrobial resistance.<sup>2</sup>

Physiological adaption to the postnatal environment in the first hours of life is critical for survival. At birth, babies transition from a semi-allogeneic, protected fetal setting to a microbe-rich extrauterine environment.<sup>3</sup> The initial mass bacterial colonisation of mucosa in the digestive, respiratory and urogenital tracts, as well as the skin, occurs during the very early neonatal period.<sup>4</sup> The maternal gut is the source of the majority of transmitted bacterial strains to the neonatal microbiome and common strains are also seeded from the maternal skin and the vaginal microbiome.<sup>5</sup> The acquisition of symbionts can positively affect gut maturation, metabolic homeostasis and immune function in early life and beyond,<sup>6</sup> but pathogenic bacteria and fungi pose an immediate risk unless contained. Adaptive immune responses require priming in neonates, and hence innate defence mechanisms play an important role in containment.

Similarly, just before delivery or during the intrapartum period babies can be infected by micro-organisms, which may lead to early-onset neonatal sepsis (EONS) especially if the mother is colonised with pathogenic bacteria.<sup>7</sup> Aspiration or ingestion of infected amniotic fluid *in utero* or infected secretions at birth are the common routes of infection.

Iron is an important commodity in the host-pathogen battle for resources.<sup>8</sup> It is a cofactor in numerous metabolic pathways that are critical for the human host as well as most pathogens. Therefore, systemic iron distribution is strictly regulated by the host.<sup>9</sup> For many human pathogens, the acquisition of iron via a variety of molecular mechanisms can

enhance growth and virulence.<sup>8</sup> Individuals with chronically high iron states (e.g. haemochromatosis) have an enhanced risk of bacterial infection,<sup>10</sup> as well as increased free radical redox damage.<sup>11</sup> Historic studies administering parenteral iron to Polynesian neonates infamously increased mortality rates by promoting *Escherichia coli* septicemias,<sup>12</sup> underlining the critical role that iron plays in the neonatal period.

Cord blood, which has high levels of serum iron and transferrin saturation,<sup>13</sup> has frequently been used as a proxy for neonatal iron status. However, previous work by our group and others has shown that healthy, vaginally-delivered, full-term babies profoundly decrease their serum iron and TSAT levels in the first 24h after birth- an adaptation that may have evolved as an innate defence against iron-requiring microorganisms.<sup>14</sup> In the current study, we hypothesised that the greater susceptibility of preterm and low birthweight babies to neonatal infections might be because they are less able to elicit this defence. We prospectively tested this in hospital-delivered Gambian neonates.

## 5.3 PARTICIPANTS AND METHODS

Full details of the methods of this study can be found in the published protocol paper.<sup>15</sup>

### Study Design

This observational study recruited well, hospital-delivered neonates into four different groups characterised according to birthweight and gestational age as shown in **Figure 1**. The primary analysis compares all babies exhibiting preterm birth (PTB), low birthweight (LBW) or both (PTB/LBW) versus full-term (FTB), normal birthweight (NBW) newborns (FTB/NBW)). Secondary analyses examine the PTB+NBW, FTB+LBW and PTB+LBW groups separately.



## **Ethics, standards and informed consent**

The study was approved by the Medical Research Council Unit The Gambia at London School of Hygiene and Tropical Medicine (MRCG at LSHTM) Scientific Coordinating Committee, the Joint Gambia Government/MRCG Ethics Committee (SCC1525) and the London School of Hygiene and Tropical Medicine Ethics Committee (Ref:14316). The study was conducted according to Good Clinical Practice (GCP) standards. All participants gave written, informed consent.

## **Study setting**

Study participants were recruited from Kanifing General Hospital (formerly Serrekunda General Hospital), in the urban Kanifing region of The Gambia, West Africa.

## **Recruitment, screening and enrolment**

In total, we planned to identify 450 healthy newly born babies during delivery at the Kanifing General Hospital (KGH) Maternity Ward. After informed consent was obtained, neonates who met the inclusion criteria were enrolled. For inclusion in the study, neonates were medically stable (no birth asphyxia nor signs of sepsis as judged by the attending study physician),  $\geq 32$  weeks gestational age and weighed  $\geq 2000\text{g}$ . Preterm babies (PTB) were  $< 37$  completed weeks gestational age (assessed by New Ballard Score<sup>16</sup>). Term babies (FTB) had a gestational age  $\geq 37$  completed weeks. Low birthweight (LBW) was defined as  $< 2500\text{g}$  in line with the usual WHO definition. Neonates who weighed  $\geq 2500\text{g}$  were defined as normal birthweight (NBW). The observational period began at delivery and lasted until the end of the 7<sup>th</sup> day of life. Data collection started on the 5<sup>th</sup> July 2017 and ended on 1<sup>st</sup> February 2019.

Pregnant mothers were excluded from the study if they were below the age of 18 years, had no fetal heartbeat detected upon admission, were known to be HIV-positive, were in receipt of *Mycobacterium tuberculosis* therapy, had taken antibiotics in the last seven days, had a

blood transfusion in the last month, were suffering from severe pre-eclampsia or antepartum haemorrhage, or were in another research study.

Babies were excluded at the delivery stage for the following reasons: major congenital malformations (not including polydactylyism), blood transfusions given to mother or neonate, severe birth asphyxia (requiring resuscitation), neonates born via breech, vacuum or via caesarean section, or a birthweight <2000g (in order to avoid the additional burden of a blood draw in these vulnerable neonates as guided by the Joint Gambia Government/MRCG Ethics Committee).

After the delivery stage, babies were excluded following the detection of infection or illness (information gained from a venous bleed or review of systems). Neonates were also removed from the study protocol if any medication other than intramuscular vitamin K, tetracycline eye ointment or immunisations were given. All medications given to mothers and neonates during the study period were recorded. Mothers who delivered multiple newborns were invited to enrol one of their neonates into the study.

### **Sample collection**

Once the neonate was fully delivered, one-minute delayed cord clamping was used (following World Health Organisation (WHO) policy<sup>17</sup>). After the umbilical cord was removed and cleaned, a trained study nurse identified the umbilical arteries and umbilical vein. Blood was collected from each using separate blood draw equipment.

At 6-24h post-delivery, recruited mothers and their neonates were invited to a private consultation with the study research clinician. Further demographic data were collected, along with a complete review of systems of the mother and neonate. Anthropometric data on the newborn was also collected at this point. Neuromuscular and physical maturation of each neonate was assessed using the New Ballard Score.<sup>16</sup> Immediately after passing the

health assessment, a venous blood draw was performed on all neonates (2 ml for PTB/LBW neonates and 3.5 ml for FTB/NBW neonates).

### **Laboratory analyses**

A full haematology panel (using a Medonic M20M GP, Boule Diagnostics, Spanga, Sweden) and glucose-6-phosphate dehydrogenase deficiency test (R&D Diagnostics Limited, Papagos, Greece) were conducted on whole blood. Serum was separated and stored at -20°C prior to analysis of ferritin, iron, unsaturated iron-binding capacity (UIBC), soluble transferrin receptor (sTfR), transferrin, c-reactive protein (CRP), haptoglobin and alpha-1-acid glycoprotein (AGP) using a fully automated biochemistry analyser (Cobas Integra 400 plus, Roche Diagnostic, Switzerland). Transferrin saturation (TSAT) was calculated. Serum samples were assessed for hepcidin concentration by ELISA (hepcidin-25 (human) EIA Kit, DRG, USA) with a dynamic range of 0.135-81 ng/mL. Hepcidin reference material refined by Diepeveen *et al*<sup>18</sup> was not used due to the lack of availability at the time of conducting laboratory analyses.

In order to ensure a consistent assessment of haemolysis in all serum samples, batches of samples were thawed before entering the biochemistry analyser and visually scored by a single operator. A previously published specimen integrity chart for haemolysis was used as reference.<sup>19</sup> Samples were scored 0 (yellow 0 g/L) to 6 (dark red 8 g/L). Samples scoring ≥5 were removed from the analysis. All serum samples experienced one freeze-thaw cycle before biochemical analyses was conducted.

### **Sample size determination**

This study targeted recruitment of 50 neonates in each of the 3 groups PTB, LBW and PTB/LBW and 300 FTB/NBW newborns. Sample size calculations for the primary outcomes of change in serum iron and TSAT between cord and the first neonatal samples were based on data from a previous study<sup>14</sup> and are summarised in the published protocol paper.<sup>15</sup> All

secondary analyses are considered exploratory and were not subjected to a formal sample size calculation.

### **Statistical analysis**

The primary analysis compared responses in the PTB and/or LBW babies combined (PTB/LBW) versus FTB/NBW. Further subgroups of the PTB/LBW group (groups A-C as summarised in **Figure 1**) were also examined in the secondary analyses (see Supplementary Material). For continuous variables, baseline characteristics are presented as means ( $\pm$  SD) for normally distributed variables. All skewed data were transformed using the ladder command in STATA. The ladder command searches a subset of the ladder of powers for a transformation that converts the variable of interest into a normally distributed variable. Results were confirmed graphically by the gladder command. Categorical variables are reported as proportions. The rate of missing data was small (<5%), thus we did not impute missing data. Participant characteristics and iron status indicators were compared using 2-tailed t-tests or  $\chi^2$  tests. Multiple regression was used to explore relationships between iron status indicators, participant characteristics, inflammatory and haematology markers. The relative strength of associations was assessed using standardised coefficients, which represent the effect on the outcome variable (expressed as a fraction of a standard deviation) caused by a one standard deviation difference in the predictor variable. Changes between cord and postnatal blood samples were assessed by paired t-tests. Comparisons of continuous variables between groups A-D were produced using repeat measures one-way analysis of variance. We examined the association between covariates and the three outcomes iron, TSAT and hepcidin status using linear regression models with backward elimination for variable selection. All models were adjusted for baseline (cord blood) levels. Unless otherwise stated, all hypothesis tests were two-sided at significance level of 0.05. All analyses were performed using DataDesk (Data Description Inc., Ithaca, USA) and STATA 15 (StataCorp LLC, Texas, USA).

## 5.4 RESULTS

**Figure 2** is the CONSORT diagram summarising subject recruitment. Baseline characteristics for the 430 neonates completing the study are shown in **Table 1**. As per protocol, there was a large difference in birthweight between NBW ( $3199\pm 376\text{g}$ ) and LBW ( $2338\pm 118\text{g}$ ) newborns, with associated differences in length and head circumference. Gestational age was also lower in the PTB babies ( $35.6\pm 0.7$ ) than FTB ( $39.4\pm 1.3\text{wk}$ ). When the LBW and PTB babies were combined as PTB/LBW, their weight and gestational ages were significantly lower than the FTB/NBW newborns. Mothers of low birthweight babies were younger ( $26.8\pm 6.6\text{y}$ ) than FTB/NBW mothers ( $29.7\pm 6.9\text{y}$ ), but there were no other differences in maternal or neonatal characteristics (**Table 1 and Supplementary Table 1 & 2**).

### Neonatal hypoferrremia

**Figure 3** illustrates a profound hypoferrremia occurring in the first day after parturition. There were no babies that failed to show a hypoferrremic response and very few in whom the response was only moderate. **Figure 4** shows that the decline had already occurred by the beginning of our sampling window at 6h post-delivery. Contrary to our initial hypothesis, there was no difference in the hypoferrremic response between any of the PTB, LBW or PTB/LBW groups and the controls. For all the babies combined, serum iron decreased over 3-fold from  $23.3\pm 7.1$  to  $7.5\pm 4.5\mu\text{mol/L}$  ( $P<0.001$ ) and TSAT decreased from  $51.7\pm 17.3$  to  $15.0\pm 6.9\%$  ( $P<0.001$ ) (**Table 2**). The mean decrease in serum iron from cord to postnatal blood was remarkably consistent across all groups (between  $15.2$  and  $16.5\mu\text{mol/L}$ ), equivalent to a range from 2.9 to 3.1-fold decrease (**Table 3**). Likewise, the spread of the decreases in TSAT between groups was very tight (between 35 and 39%), equivalent a 3.4 to 3.5-fold reduction. Correspondingly there was a substantial increase in UIBC in all groups indicating an enhanced ability to sequester any free iron ( $22.9\pm 10.5$  to  $43.5\pm 15.9\mu\text{mol/L}$ ,

$P<0.0001$  for all babies combined). Notably, ferritin, TIBC and haemoglobin levels increased from cord to the first neonatal bleed, and there was no change in transferrin; thus confirming that the decrease in serum iron was an active adaptation not related to altered haemodynamics.

**Figure 5** shows that all groups had similar hepcidin levels ( $20.9\pm13.8$  in PTB/LBW vs  $19.4\pm14.4$  ng/mL in FTB/NBW,  $P=0.3$ ) in their cord blood. However, the cord blood from PTB neonates had slightly higher TSAT values ( $54.8\pm17.7$  vs  $50.2\pm16.9\%$ ,  $P=0.017$ ) and serum iron ( $24.4\pm7.3$  vs  $22.7\pm7.0$   $\mu\text{mol/L}$ ,  $P=0.01$ ) in comparison to FTB neonates. In the postnatal venous blood samples, all groups had similar hepcidin levels ( $37.4\pm23.5$  in PTB/LBW vs  $38.9\pm23.9$  ng/ml in FTB/NBW,  $P=0.5$ ). The venous blood from PTB and LBW neonates had slightly higher TSAT values ( $16.1\pm8.4\%$  and  $16.2\pm5.3\%$  vs  $14.4\pm6.0\%$ , both  $P=0.01$ ) in comparison to FTB neonates. There is no evidence that these small differences in TSAT will translate into a clinically important difference in susceptibility to neonatal sepsis or infection. Additional comparisons can be seen in **Supplementary Tables 3 & 4**.

Comparisons of further iron, infection and haematological parameters in umbilical cord and venous blood can be seen in **Supplementary Table 5**. Analysis of the babies subdivided into groups A-D is listed in **Supplementary Tables 6, 7 and 8**. As anticipated, based upon the lack of difference in hypoferrremia between PTB, LBW and FTB/NBW neonates, there were no differences between the additional subgroupings of A-D.

### **Factors associated with the decline in serum iron and TSAT**

CRP levels increased by over 10-fold between cord and postnatal blood ( $0.17\pm0.6$  to  $2.16\pm4.0$  mg/L,  $P<0.001$ ) and hepcidin levels doubled ( $19.9\pm14.2$  to  $38.4\pm23.7$  ng/ml,  $P<0.001$ ). Figure 4 illustrates the timecourse of the changes.

**Table 3** lists the results of the regression analysis of factors associated with Day 1 serum iron. Use of standardised coefficients permits ranking in order of the effect size per standard deviation of the predictor variable. In univariate analysis CRP, hepcidin and transferrin were most strongly associated with serum iron. Note that CRP was entered as the reciprocal of the square root, so the direction of the coefficient is reversed (i.e. a high CRP was associated with a low serum iron). Haptoglobin, age of mother and birthweight were also significantly associated with serum iron. In multivariable analysis, CRP and hepcidin were the strongest correlates. The parallel analysis for TSAT showed broadly similar associations (**Table 3**).

#### **Factors associated with postnatal hepcidin levels**

Based on the *pre-hoc* assumption that hepcidin orchestrates the postnatal hypoferremia, we examined the factors associated with neonatal hepcidin levels on Day 1 (**Table 4**). The strongest predictor was time of bleed with a negative coefficient. Examination of **Figure 4C** suggests that this was because hepcidin rose very fast at, or immediately after, parturition and was already starting to decline by 6h. Ferritin was positively associated with hepcidin, and serum iron was negatively correlated. Surprisingly there was no evidence of a cross-sectional association between hepcidin and CRP.

## **5.5 DISCUSSION**

The rapid and profound postnatal hypoferremia demonstrated in this study closely matches our prior findings in full-term, vaginally-delivered rural Gambian babies<sup>14</sup> and the results of studies elsewhere.<sup>20,21</sup> The 3-4-fold decrease recorded here is towards the top end of the 2-4-fold range recorded in prior studies.<sup>14,20,21</sup> We have previously proposed that this may represent an evolved innate immune response designed to deprive blood-stream bacteria of iron and hence create a hostile bacteriostatic environment.<sup>14</sup> The current study was

designed to test the hypothesis that immature or growth-restricted neonates might have a lesser ability to trigger this innate defence and that this might explain their greater susceptibility to septicemias.<sup>22</sup> Our hypothesis was firmly refuted. We showed that the premature and low birthweight neonates all exhibited a profound hypoferremia during the first 24h of life, with no detectable differences from the full-term, normal birthweight newborns. In fact, there was a remarkable similarity in the hypoferremic response across all the study groups that underscores the efficiency of the process and supports the likelihood that it occurs by the process of evolution.

The host-pathogen battle for iron has been extensively studied<sup>23</sup> and it has long been assumed that the hypoferremia of the acute phase response acts as an innate defence against iron-requiring organisms. Hepcidin is a key, though not necessarily the only,<sup>24</sup> regulator of this response.<sup>25</sup> Inflammatory cytokines including IL-6, IL-22 and Type-1 interferon rapidly upregulate hepcidin expression and release from the liver.<sup>26</sup> Hepcidin causes hypoferremia by inhibiting the action of the transmembrane iron-exporter ferroportin in enterocytes and macrophages, thus blocking iron absorption and recycling.<sup>27</sup> Injection of recombinant hepcidin results in the very rapid induction of hypoferremia in mice<sup>28</sup> and humans.<sup>29</sup>

In both univariate and multivariate analysis neonatal serum iron and TSAT levels were most strongly correlated with CRP and hepcidin (**Table 3**) suggesting that the hypoferremia is, at least partly, driven by an inflammatory response to the stress of the birth process and/or the exposure of the newborn to vaginal and/or gastrointestinal micro-organisms not previously encountered *in utero*. The modest proportion of variance explained in the full multivariate model (29% for serum iron and 22% for TSAT) might reflect the fact that both measures have been suppressed too close to their physiological lower limit and hence display a limited range. It is also possible that hepcidin-independent mechanisms, driven by effector molecules unmeasured in this study, are playing an additional role.<sup>24</sup> A full examination of



the mechanism responsible for the hypoferremia would probably require studies in an animal model. Neonatal piglets may be a suitable model due to their low iron endowment at birth, low iron milk<sup>30</sup>, fast growth rate<sup>31</sup> and similarities to human anatomy and physiology.<sup>32</sup> These factors result in young porcine models often becoming iron deficient and anaemic without the need for low-iron diets (i.e. as seen in murine models).<sup>33</sup>

Using *ex vivo* assays with sentinel bacteria, we have previously demonstrated that low TSAT values in serum from adults<sup>34</sup> and neonates<sup>14</sup> exerts a powerful bacteriostatic effect. In the neonatal study, growth rates of *Escherichia coli*, *Streptococcus pneumoniae*, *Streptococcus agalactiae* and *Staphylococcus aureus*, were highly significantly lower in neonatal serum than in cord serum and for each organism growth rates were significantly associated with TSAT. *S. aureus* was least responsive, possibly reflecting its ability to utilise haem iron, though it was still clearly influenced by transferrin saturation. *E. coli* was most responsive, which may explain why intramuscular iron administration to Polynesian neonates increased septicaemia rates with a major shift towards *E. coli* as the most frequently identified cause.<sup>12</sup> The hypoferremia observed in the current hospital-based study (in both PTB/LBW and FTB/NBW babies) was more profound than we previously observed in rural home deliveries<sup>14</sup> (TSAT declined to 15% versus 24% in rural babies) and hence the bacteriostatic effect would be expected to be even greater.

There are several strengths and limitations to our study. The large sample size allows us to confidently exclude any clinically-meaningful differences in the postnatal response of preterm and low birthweight babies compared to the full-term normal birthweight neonates. By protocol, we did not recruit mothers with complex medical histories (e.g. pre-eclampsia, antepartum haemorrhage or antenatal infection) or sick babies (e.g. with birth asphyxia or suspected sepsis). We did not recruit newborns born <32 completed weeks gestation and/or <2000g birthweight, or those delivered via c-section, vacuum or forceps. We cannot speculate whether similar hypoferremic responses would occur in such cases, but

reciprocally we can conclude that the responses we observed are a characteristic feature of normal human birth and were not elicited by pathological circumstances. In order to allow mothers to recover from their delivery and the neonates to be stabilised and checked for inclusion, we constrained our first postnatal blood draw to after 6h post-delivery. With many samples collected soon after this point, we have demonstrated that the reduction in circulating iron levels occurs very rapidly, but we cannot state how rapidly. We did not measure pro-inflammatory cytokines (e.g. IL-6 and IL-22) which might have provided additional insights into the mechanisms eliciting hypoferremia. Similarly, we did not attempt to address any possible hepcidin-independent mechanisms because the putative mediators remain unknown. Such studies may require animal models.

In conclusion, our results suggest that the innate postnatal iron restriction strategy in the first hours of life has evolved as an intrinsic mechanism to protect neonates from common pathogens and/or free-radical damage, and occurs regardless of gestational age or birthweight. This hypoferremia is at least partly mediated by the hormone, hepcidin. The trigger mechanism, relation to maternal iron status, and its effect on susceptibility to systemic bacterial infections require further investigation.

Our results highlight the importance of hypoferremia as a conserved mechanism of protection, and prompt further research into the use of iron restriction as a transient bacteriostatic mechanism to limit bacterial growth and virulence in other instances of infection. Hypoferremia can slow the multiplication of bacterial pathogens,<sup>34</sup> which in combination with antibiotics, could allow enough time for the adaptive immune system to fight the infection. Augmentation of innate immunity in neonates and other at-risk groups (elderly or immunocompromised) might be achieved by the use of small molecule orally-administered mini-hepcidins currently under development as hepcidin agonists.<sup>29,35</sup>

## **Abbreviations**

AGP = Alpha-1-acid glycoprotein

CRP = C-reactive protein

ELISA = Enzyme-linked immunosorbent assay

EONS = Early-onset neonatal sepsis

FTB = Full-term birth

G6PD = Glucose-6-phosphate dehydrogenase

GCP = Good Clinical Practice

IL-6 = Interleukin 6

IL-22 = Interleukin 22

KGH = Kanifing General Hospital

KMC = Kanifing Municipal Council

LBW = Low birthweight

MRCG = Medical Research Council Unit The Gambia at LSHTM

NBW = Normal birthweight

PI = Principal investigator

PTB = Preterm birth

SCC = Scientific Coordinating Committee

sTfR = Soluble transferrin receptor

TIBC = Total iron-binding capacity

TLR = Toll-like receptor

TSAT = Transferrin saturation

UIBC = Unbound iron-binding capacity

V1 = Venous bleed 1

WHO = World Health Organisation

**Acknowledgements:**

We thank Kanifing Municipal Council (KMC) and Kanifing General Hospital (KGH) for their support during the study.

**Funding:**

The research was undertaken with a research grant provided by the Bill & Melinda Gates Foundation (OPP1152353). The funding agency had no role in the design and conduct of the study, and did not have any in the collection, management, analyses or interpretation of the data nor in the preparation, review, or approval of the manuscript. The Nutrition Theme of the MRC Unit The Gambia at LSHTM are supported by core funding MC-A760-5QX00 to the MRC Unit The Gambia/MRC International Nutrition Group by the UK MRC and the UK Department for the International Development (DFID) under the MRC/DFID Concordat agreement.

**Availability of Data and Materials:**

All data will be made available to researchers upon reasonable request to the study PI and clearance by the MRCG Scientific Coordinating and Ethics Committees.

**Authors' Contributions:**

**JHC** is PhD student with MRCG at LSHTM. He contributed to the development of the protocols, study management, field data collection, development of the data analysis plan, conducting the data analysis and drafting of the manuscript.

**OJ** is a research clinician with MRCG at LSHTM. He contributed to the development of the protocols, study management, field data collection and patient care.

**NIM** is a statistician at MRCG at LSHTM. He wrote the data analysis plan in consultation with JC, CC and AMP.

**SRG** was a visiting B.Sc. student from Cardiff University. He contributed to the laboratory analyses and field data collection.

**BJBT** is a laboratory technician with MRCG at LSHTM. He contributed to the laboratory analyses and field data collection.

**AMP** is a Professor of International Nutrition at LSHTM and Nutrition Theme Leader at MRCG at LSHTM. He conceived the study, obtained the funding, contributed to the development of the protocols, the data analysis plan, conducting the data analysis and the drafting of the manuscript.

**CC** is Senior Investigator Scientist at MRCG at LSHTM and Group Leader for Iron, Infection and Anaemia. She was the PI on the trial, conceived the study, and obtained the funding. She was responsible for the overall development of the protocols, the data analysis plan, conducting the data analysis and the drafting of the manuscript.

All authors reviewed the final manuscript prior to submission.

**Competing Interests:**

The authors declare that they have no competing interests.

**Consent for Publication:**

Not applicable.

**Ethics Approval and Consent to Participate:**

This study has been approved by The Gambia Government/MRC Joint Ethics Committee (no. SCC1525) and London School of Hygiene and Tropical Medicine Ethics Committee (ref no. 14316). The study was conducted according to Good Clinical Practice (GCP) standards. The study procedures were explained to the neonate's mother/guardians orally and in writing. A neonate was only recruited into the study after the written, informed consent was provided by the mother/guardian.

## Author Details:

<sup>1</sup>Medical Research Council Unit The Gambia at the London School of Hygiene & Tropical Medicine, Atlantic Boulevard, Fajara, P.O. Box 273, Banjul, The Gambia

<sup>2</sup>Cardiff University, Cardiff, Wales, CF10 3AT, United Kingdom

## 5.6 REFERENCES

- 1 GBD 2016 Causes of Death Collaborators M, Abajobir AA, Abbafati C, *et al.* Global, regional, and national age-sex specific mortality for 264 causes of death, 1980-2016: a systematic analysis for the Global Burden of Disease Study 2016. *Lancet* 2017; **390**: 1151–210.
- 2 Laxminarayan R, Duse A, Wattal C, *et al.* Antibiotic resistance—the need for global solutions. *Lancet Infect Dis* 2013; **13**: 1057–98.
- 3 Moore RE, Townsend SD. Temporal development of the infant gut microbiome. *Open Biol* 2019; **9**: 190128.
- 4 Dominguez-Bello MG, Costello EK, Contreras M, *et al.* Delivery mode shapes the acquisition and structure of the initial microbiota across multiple body habitats in newborns. *Proc Natl Acad Sci* 2010; **107**: 11971–5.
- 5 Ferretti P, Pasolli E, Tett A, Huttenhower C, Correspondence NS. Mother-to-Infant Microbial Transmission from Different Body Sites Shapes the Developing Infant Gut Microbiome. *Cell Host Microbe* 2018; **24**: 133–45.
- 6 Dominguez-Bello MG, Godoy-Vitorino F, Knight R, Blaser MJ. Role of the microbiome in human development. *Gut* 2019; **68**: 1108–14.
- 7 Chan GJ, Lee AC, Baqui AH, Tan J, Black RE. Prevalence of early-onset neonatal infection among newborns of mothers with bacterial infection or colonization: a systematic review and meta-analysis. *BMC Infect Dis* 2015; **15**: 118.
- 8 Schaible UE, Kaufmann SHE. Iron and microbial infection. *Nat Rev Microbiol* 2004; **2**:

- 946–53.
- 9 Pigeon C, Ilyin G, Courselaud B, *et al.* A New Mouse Liver-specific Gene, Encoding a Protein Homologous to Human Antimicrobial Peptide Heparin, Is Overexpressed during Iron Overload. *J Biol Chem* 2001; **276**: 7811–9.
  - 10 Khan FA, Fisher MA, Khakoo RA. Association of hemochromatosis with infectious diseases: expanding spectrum. *Int J Infect Dis* 2007; **11**: 482–7.
  - 11 Pietrangolo A. Hereditary hemochromatosis. *Biochim Biophys Acta - Mol Cell Res* 2006; **1763**: 700–10.
  - 12 Barry DM, Reeve AW. Increased incidence of gram negative neonatal sepsis with intramuscular iron administration. *Pediatrics* 1977; **60**: 908-12.
  - 13 Lorenz L, Peter A, Poets CF, Franz AR. A Review of Cord Blood Concentrations of Iron Status Parameters to Define Reference Ranges for Preterm Infants. *Neonatology* 2013; **104**: 194–202.
  - 14 Prentice S, Jallow AT, Sinjanka E, *et al.* Heparin mediates hypoferrremia and reduces the growth potential of bacteria in the immediate post-natal period in human neonates. *Sci Rep* 2019; **9**: 16596.
  - 15 Cross JH, Jarjou O, Mohammed NI, Prentice AM, Cerami C. Neonatal iron distribution and infection susceptibility in full term, preterm and low birthweight babies in urban Gambia: study protocol for an observational study. *Gates Open Res* 2019; **3**: 1469.
  - 16 Ballard JL, Khoury JC, Wedig K, Wang L, Eilers-Walsman BL, Lipp R. New Ballard Score, expanded to include extremely premature infants. *J Pediatr* 1991; **119**: 417–23.
  - 17 WHO | Delayed umbilical cord clamping for improved maternal and infant health and nutrition outcomes. WHO. 2019. Available at: [https://www.who.int/nutrition/publications/guidelines/cord\\_clamping/en/](https://www.who.int/nutrition/publications/guidelines/cord_clamping/en/).
  - 18 Diepeveen LE, Laarakkers CMM, Martos G, *et al.* Provisional standardization of heparin assays: creating a traceability chain with a primary reference material, candidate reference method and a commutable secondary reference material. *Clin*

- Chem Lab Med* 2019; **57**: 864–72.
- 19 Dugan L, Leech L, Speroni KG, Corriher J. Factors Affecting Hemolysis Rates in Blood Samples Drawn From Newly Placed IV Sites in the Emergency Department. *J Emerg Nurs* 2005; **31**: 338–45.
  - 20 Szabo M, Vasarhelyi B, Balla G, *et al.* Acute postnatal increase of extracellular antioxidant defence of neonates: the role of iron metabolism. *Acta Paediatr* 2001; **90**: 1167–70.
  - 21 Balogh Á, Szabó M, Kelen D, *et al.* Prohepcidin levels during human perinatal adaptation. *Pediatr Hematol Oncol* 2007; **24**: 361-8
  - 22 Schrag SJ, Farley MM, Petit S, *et al.* Epidemiology of invasive early-onset neonatal sepsis, 2005 to 2014. *Pediatrics*. 2016; **138**: e20163013.
  - 23 Skaar EP. The battle for iron between bacterial pathogens and their vertebrate hosts. *PLoS Pathog* 2010; **6**: e1000949.
  - 24 Enculescu M, Metzendorf C, Sparla R, *et al.* Modelling Systemic Iron Regulation during Dietary Iron Overload and Acute Inflammation: Role of Heparin-Independent Mechanisms. *PLoS Comput Biol* 2017; **13**: e1005322.
  - 25 Michels K, Nemeth E, Ganz T, Mehrad B. Heparin and Host Defense against Infectious Diseases. *PLoS Pathog*. 2015; **11**: e1004998.
  - 26 Ganz T. Systemic iron homeostasis. *Physiol Rev* 2013; **93**: 1721–41.
  - 27 Nemeth E, Tuttle MS, Powelson J, *et al.* Heparin regulates cellular iron efflux by binding to ferroportin and inducing its internalization. *Science* 2004; **306**: 2090–3.
  - 28 Rivera S, Nemeth E, Gabayan V, Lopez MA, Farshidi D, Ganz T. Synthetic hepcidin causes rapid dose-dependent hypoferremia and is concentrated in ferroportin-containing organs. *Blood* 2005; **106**: 2196–9.
  - 29 Protagonist Therapeutics. Protagonist Therapeutics announces final phase 1 study results with novel hepcidin mimetic, PTG-300. <https://www.prnewswire.com/news-releases/protagonist-therapeutics-announces-final-phase-1-study-results-with-novel-hepcidin-mimetic-ptg-300-300571268.html>. (accessed Sept 18, 2019).



- 30 Starzyński RR, Laarakkers CMM, Tjalsma H, *et al.* Iron Supplementation in Suckling Piglets: How to Correct Iron Deficiency Anemia without Affecting Plasma Hepcidin Levels. *PLoS One* 2013; **8**: e64022.
- 31 Ventrella D, Dondi F, Barone F, *et al.* The biomedical piglet: establishing reference intervals for haematology and clinical chemistry parameters of two age groups with and without iron supplementation. *BMC Vet Res* 2016; **13**: 23.
- 32 Moughan PJ, Birtles MJ, Cranwell PD, Smith WC, Pedraza M. The piglet as a model animal for studying aspects of digestion and absorption in milk-fed human infants. *World Rev. Nutr. Diet.* 1992; **67**: 40-113
- 33 Li Y, Hansen SL, Borst LB, Spears JW, Moeser AJ. Dietary Iron Deficiency and Oversupplementation Increase Intestinal Permeability, Ion Transport, and Inflammation in Pigs. *J Nutr* 2016; **146**: 1499-505.
- 34 Cross JH, Bradbury RS, Fulford AJ, *et al.* Oral iron acutely elevates bacterial growth in human serum. *Sci Rep* 2015; **5**: 16670.
- 35 Vifor Pharma. Vifor Pharma ferroportin inhibitor enters phase-I clinical trial. <http://www.viforpharma.com/en/media/press-releases/201802/2167159> (accessed Sept 18, 2019).

## Figure and Table Legends:

### Figure 1: Schematic diagram of all study groupings and the generation of subgroups.

PTB = preterm birth, FTB = full-term birth, LBW = low birthweight and NBW = normal birthweight. FTB/NBW neonates are FTB+NBW. PTB/LBW neonates are FTB+LBW, PTB+NBW and PTB+LBW neonates.

### Figure 2: CONSORT diagram for participant flow. PTB/LBW are displayed in GREEN, and FTB/NBW in ORANGE.

### Figure 3: Analysis of serum iron (A), TSAT (B) and hepcidin (C) in umbilical cord (BLUE) and post-natal venous blood (RED) based on all study groupings. Horizontal lines represent the arithmetic group means. FTB/NBW are FTB+NBW. PTB/LBW are FTB+LBW, PTB+NBW and PTB+LBW neonates. \*\*\* = all group comparisons between cord and venous blood are statistically significant ( $P < 0.001$ ). Hepcidin displayed as $\log_{10}$ .

### Figure 4: Timecourse of the changes in serum iron (A), TSAT (B), hepcidin (C) and CRP (D) in the first day of life. Means $\pm$ 95% CI. PTB/LBW are RED columns and significance lines. FTB/NBW are BLUE columns and significance lines. Columns are plotted according to mean time of bleed for the categories 0, 1-8, 9-16 and 17-24 hours. \*\*\*\* = $P < 0.0001$ , \*\*\* = $P < 0.001$ , \*\* = $P < 0.01$ , \* = $P < 0.05$ . No significance line = $P > 0.05$ .

### Figure 5: Comparisons of serum iron (A = cord, B = venous), TSAT (C = cord, D = venous) and hepcidin (E = cord, F = venous) in cord and post-natal venous blood. Means $\pm$ 95% CI. FTB/NBW are BLUE columns. PTB/LBW are RED columns. PTB are DARK GREY columns. LBW are LIGHT GREY columns. Significance lines represent the comparison of PTB/LBW, PTB or LBW groups to the FTB/NBW group. \*\* = $P < 0.01$ , \* = $P < 0.05$ . No significance line = $P > 0.05$ .

**Table 1: Participant Characteristics.** Demographic, clinical and pregnancy outcome characteristics of the women and their newborns. Data are presented as arithmetic mean ( $\pm$  SD) and analysed by one-way analysis of variance or as a proportion (%) and analysed by Pearson  $\chi^2$  test. *P* values in bold font are considered significant based on  $P < 0.05$ .

**Table 2: Iron Parameters (cord vs venous sample) for FTB/NBW, PTB/LBW, PTB and LBW neonates.** Data are presented as mean ( $\pm$  SD) and analysed by one-way analysis of variance. *P* values in bold font are considered significant based on  $P < 0.05$ . Skewed variables were transformed as follows:  $\log_{10}$  (serum iron),  $1/\sqrt{\text{TIBC}}$ ,  $\sqrt{\text{ferritin}}$ ,  $1/\sqrt{\text{CRP}}$ ,  $\log_{10}$  (hepcidin).

**Table 3: Linear regression of factors associated with postnatal serum iron and TSAT.** All regressions were adjusted for 4 grades of visually assessed haemolysis. Ranked according to their standardized coefficients calculated in STATA. <sup>a</sup>  $1/\sqrt{\text{CRP}}$ , <sup>b</sup>  $\log_{10}$  (hepcidin).

**Table 4: Linear regression of factors associated with postnatal serum hepcidin.** All regressions were adjusted for 4 grades of visually assessed haemolysis. Ranked according to their standardized regression coefficients calculated in STATA. <sup>a</sup>  $\sqrt{\text{ferritin}}$ , <sup>b</sup>  $\log_{10}$  (serum iron), <sup>c</sup>  $\sqrt{\text{sTfR}}$ .

Figure 1

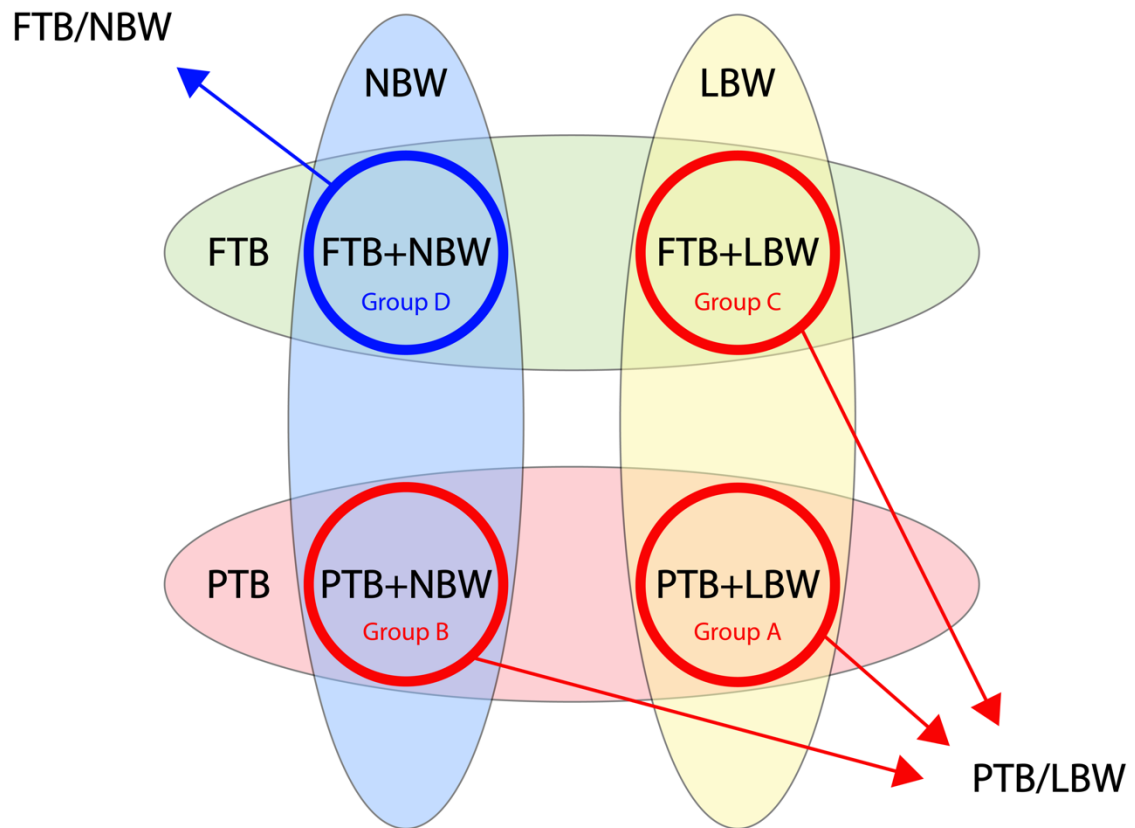


Figure 2

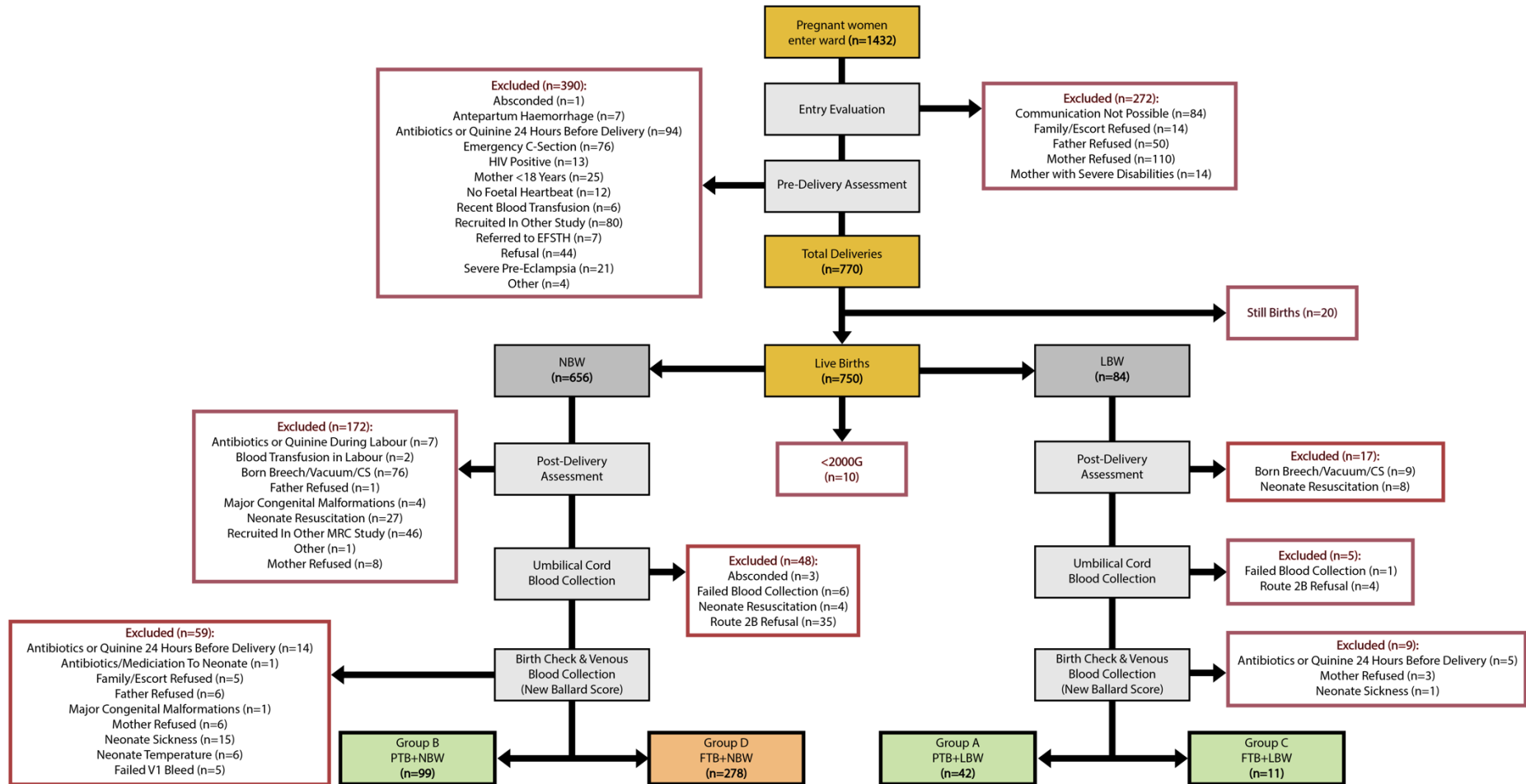


Figure 3

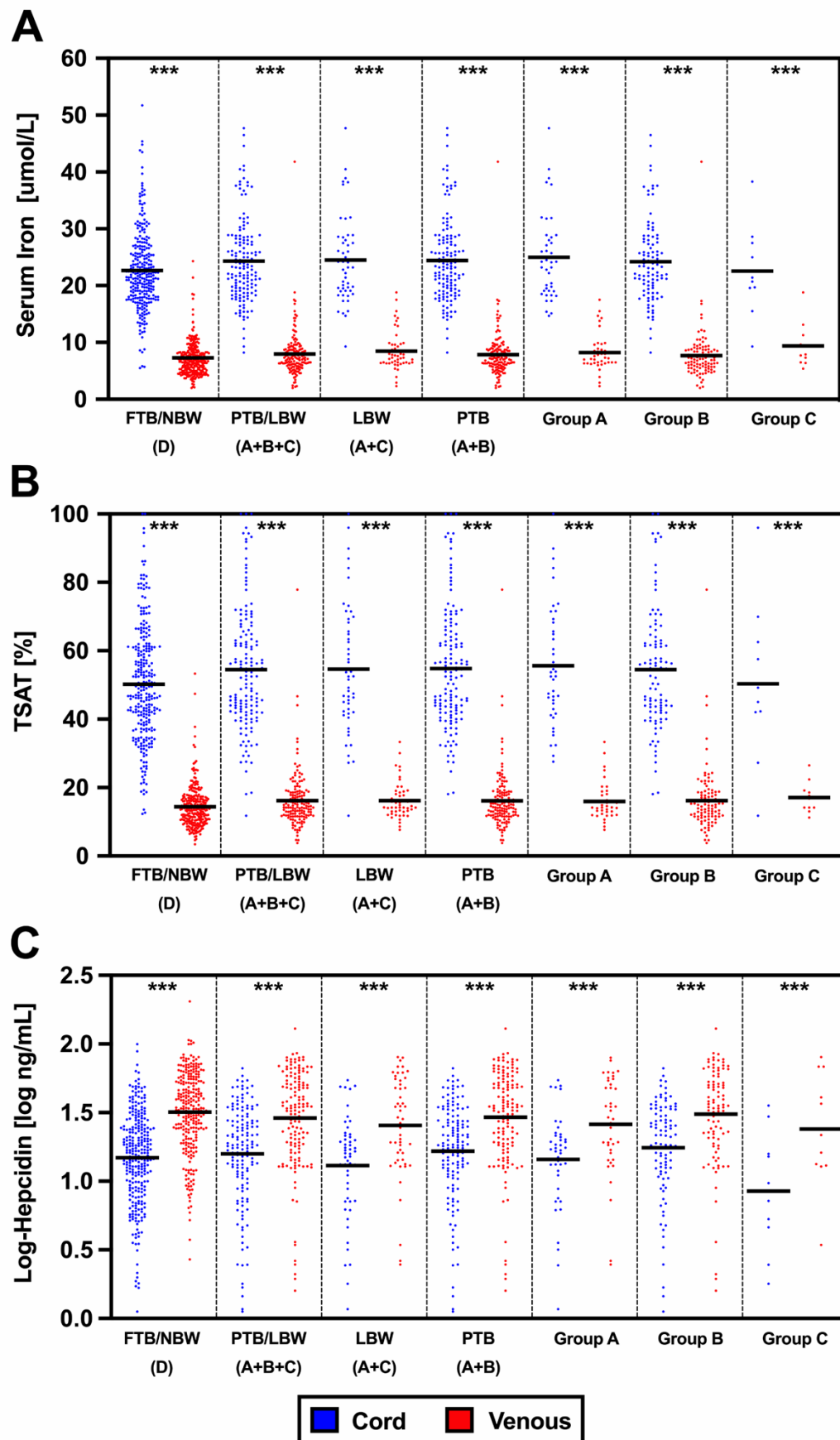
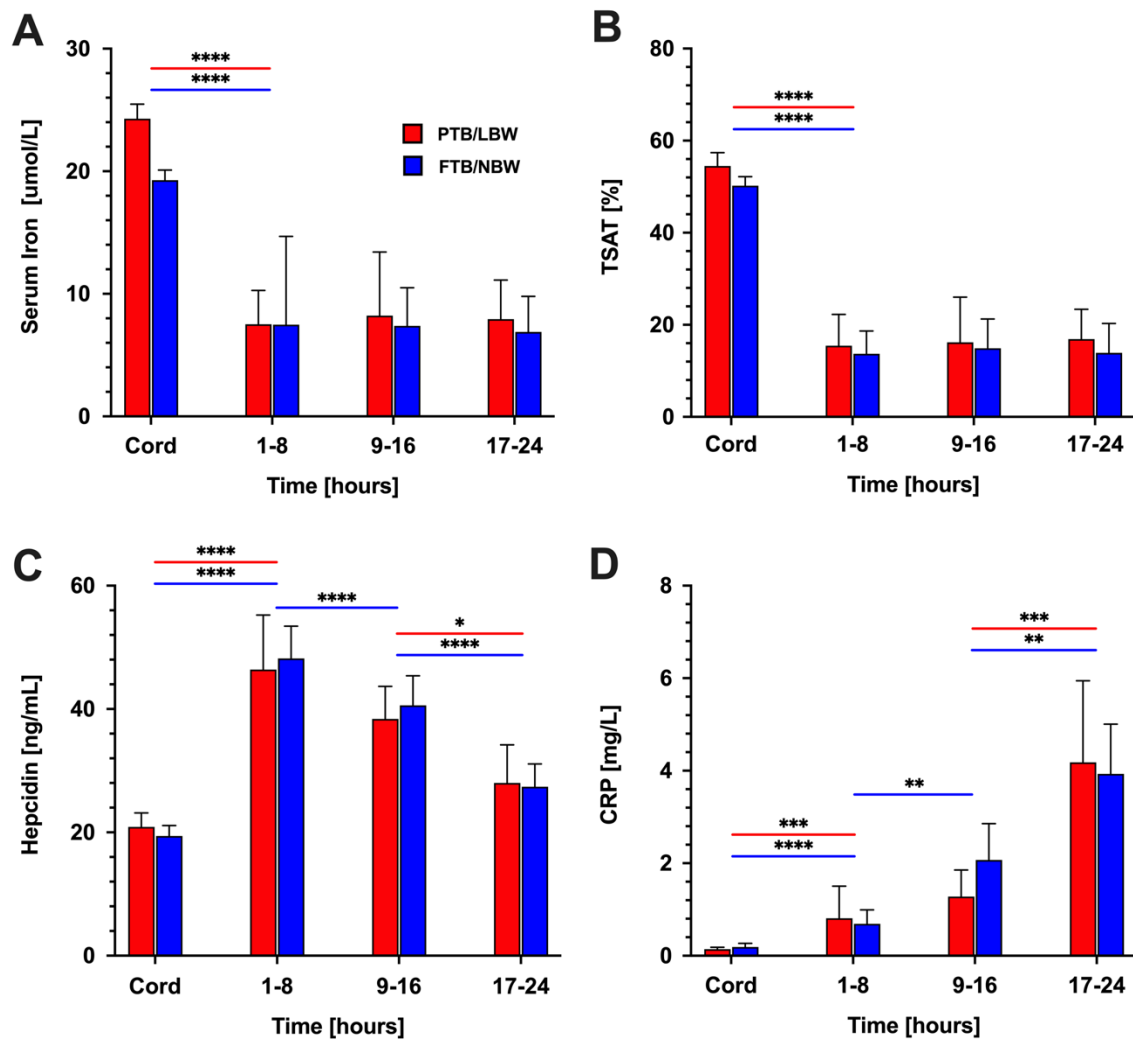
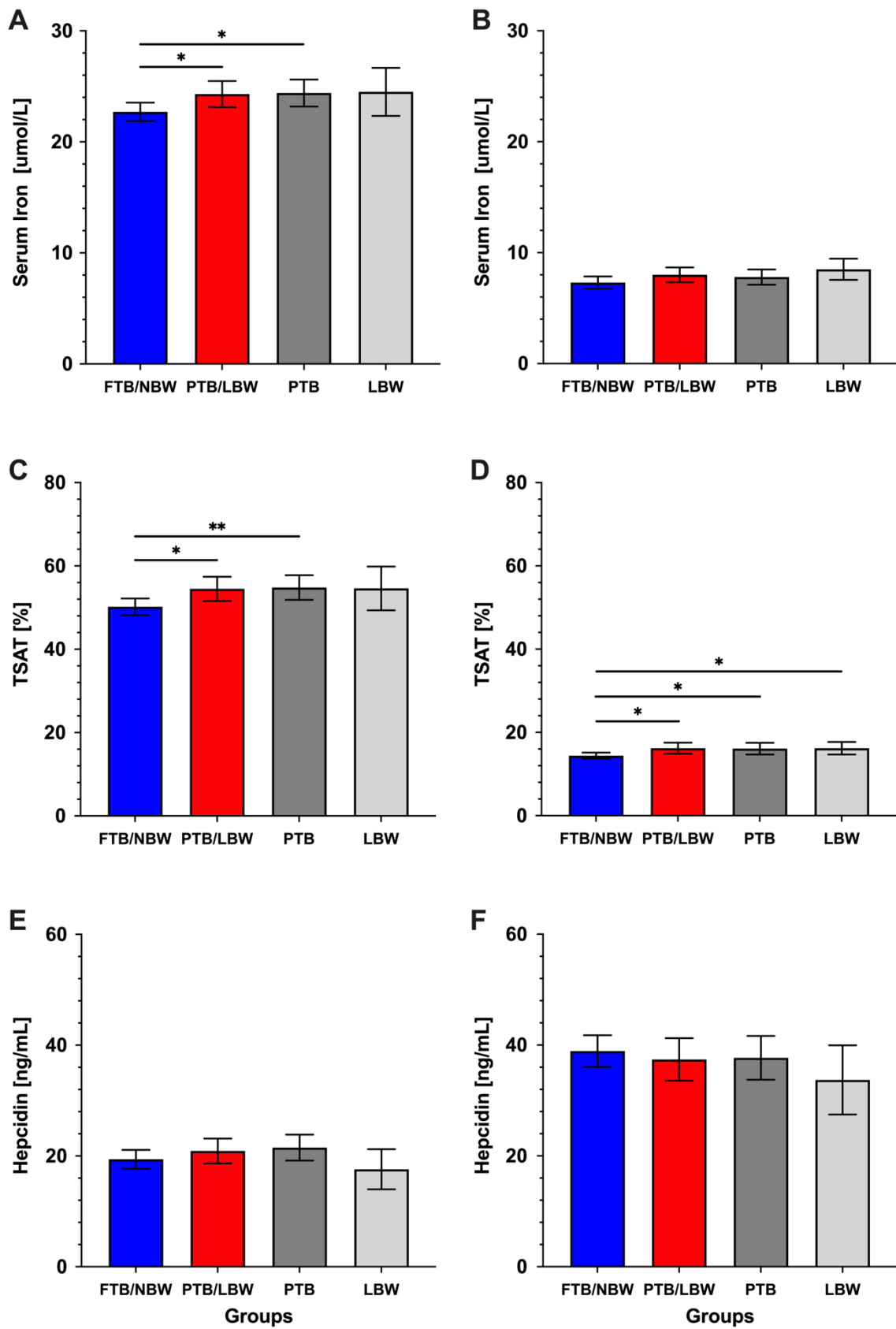


Figure 4



**Figure 5**





**Table 1 – Participant characteristics**

	Whole Population	FTB/NBW	PTB/LBW	<i>P value</i>	FTB	PTB	<i>P value</i>	NBW	LBW	<i>P value</i>
Number of participants (n)	430	278	152	..	289	141	..	377	53	..
Characteristic	Mean (± SD)									
Gestational age (Weeks)	38.1 (±2.1)	39.4 (±1.3)	35.8 (±1.1)	<0.001	39.4 (±1.3)	35.6 (±0.7)	<0.001	38.4 (±2.0)	35.9 (±1.7)	<0.001
Birth weight (g)	3092 (±453)	3299 (±368.3)	2715 (±338.5)	<0.001	3264 (±402.2)	2741 (±336.9)	<0.001	3199 (±376.3)	2338 (±117.7)	<0.001
Head circumference (cm)	34.0 (±1.2)	34.6 (±1.0)	33.0 (±0.1)	<0.001	34.5 (±1.1)	33.1 (±0.1)	<0.001	34.3 (±1.1)	32.3 (±0.8)	<0.001
Length (cm)	49.2 (±1.9)	50.0 (±1.6)	47.7 (±1.6)	<0.001	49.9 (±1.7)	47.7 (±1.6)	<0.001	49.6 (±1.7)	46.7 (±1.4)	<0.001
Maternal haemoglobin ≤7 days before delivery (g/dl)	11.6 (±1.8)	11.6 (±1.8)	11.4 (±1.8)	0.25	11.6 (±1.8)	11.5 (±1.8)	0.35	11.5 (±1.8)	11.8 (±1.7)	0.75
Age of mother (Years)	29.4 (±6.9)	29.7 (±6.9)	28.7 (±6.8)	0.16	29.6 (±7.0)	28.9 (±6.8)	0.35	29.7 (±6.9)	26.8 (±6.6)	0.003
1 min APGAR score (0-10)	9.7 (±0.7)	9.6 (±0.8)	9.7 (±0.6)	0.51	9.6 (±0.8)	9.8 (±0.6)	0.26	9.7 (±0.7)	9.6 (±0.8)	0.7
Time from admission to birth (Hours)	3.2 (±5.1)	3.2 (±5.2)	3.2 (±4.8)	0.96	3.2 (±5.2)	3.2 (±4.8)	0.94	3.2 (±5.1)	3.3 (±5.0)	0.94
Time from delivery to postnatal blood collection (Hours)	12.7 (±5.3)	12.7 (±5.4)	12.8 (±5.3)	0.81	12.6 (±5.3)	13.0 (±5.3)	0.47	12.8 (±5.3)	12.5 (±4.8)	0.71
	Percentage (%)									
Percentage male (%)	53.5% (224)	54.3% (151)	52.0% (79)	0.64	54.7% (158)	51.1% (72)	0.48	53.8% (203)	50.9% (27)	0.69
Percentage early term, ≥37-≤38 weeks (%)	19.3% (83)	27.7% (77)	3.9% (6)	<0.001	28.7% (83)	0.0% (0)	<0.001	20.4% (77)	11.3% (6)	0.12
G6PD deficiency positive (%)	12.6% (54)	11.5% (32)	14.6% (22)	0.36	11.4% (33)	15.0% (21)	0.3	13.3% (50)	7.7% (4)	0.26
Multiple births (%)	3% (13)	2.2% (6)	4.6% (7)	0.16	2.1% (6)	4.9% (7)	0.1	2.7% (10)	5.7% (3)	0.23
Percentage of mothers on antenatal iron/folic acid (%)	82.1% (353)	81.7% (227)	82.9% (126)	0.87	81.0% (234)	84.4% (119)	0.83	83.3% (314)	73.6% (39)	0.1

**Table 2 - Changes in iron status, inflammation and hepcidin between cord and postnatal blood**

Sample Type	Whole Population					FTB/NBW					PTB/LBW					Preterm Birth (PTB)					Low Birthweight (LBW)				
	n	Cord	n	Venous	P value	n	Cord	n	Venous	P value	n	Cord	n	Venous	P value	n	Cord	n	Venous	P value	n	Cord	n	Venous	P value
Serum iron (µmol/L)	425	23.3 (±7.1)	421	7.5 (±4.5)	<0.001	275	22.7 (±7.0)	271	7.3 (±4.6)	<0.001	150	24.3 (±7.3)	150	8.0 (±4.1)	<0.001	140	24.4 (±7.3)	139	7.8 (±4.1)	<0.001	51	24.5 (±7.7)	51	8.5 (±3.4)	<0.001
UIBC (µmol/L)	423	22.9 (±10.5)	420	43.5 (±15.9)	<0.001	273	23.7 (±10.4)	271	44.1 (±18.0)	<0.001	150	21.5 (±10.6)	149	42.2 (±10.9)	<0.001	140	21.2 (±9.8)	138	42.0 (±10.9)	<0.001	51	22.1 (±12.2)	50	43.8 (±10.2)	<0.001
TIBC (µmol/L)	423	46.1 (±8.0)	420	51.0 (±17.9)	<0.001	273	46.3 (±8.1)	271	51.4 (±20.7)	<0.001	150	45.8 (±7.8)	149	50.2 (±11.3)	<0.001	140	45.6 (±7.4)	138	49.8 (±11.1)	<0.001	51	46.6 (±9.3)	50	52.3 (±11.9)	<0.001
Transferrin (g/L)	426	1.96 (±0.3)	423	1.99 (±0.3)	0.003	275	1.98 (±0.3)	273	1.98 (±0.3)	0.8335	151	1.93 (±0.3)	150	2.00 (±0.3)	<0.001	141	1.92 (±0.3)	139	1.99 (±0.3)	<0.001	52	1.94 (±0.4)	51	2.02 (±0.4)	0.0024
TSAT (%)	423	51.7 (±17.3)	420	15.0 (±6.9)	<0.001	273	50.2 (±16.7)	271	14.4 (±6.1)	<0.001	150	54.5 (±18.0)	149	16.2 (±8.2)	<0.001	140	54.8 (±17.7)	138	16.1 (±8.4)	<0.001	51	54.6 (±18.7)	50	16.2 (±5.3)	<0.001
Serum ferritin (ug/L)	426	210.5 (±157.8)	415	381.2 (±297.6)	<0.001	275	212.6 (±157.7)	271	393.9 (±312.6)	<0.001	151	206.6 (±158.5)	149	358.6 (±268.1)	<0.001	141	206.8 (±154.8)	138	355.7 (±259.6)	<0.001	52	192.1 (±179.5)	51	366.2 (±343.8)	<0.001
Serum AGP (g/L)	426	0.18 (±0.1)	423	0.24 (±0.2)	<0.001	275	0.19 (±0.1)	273	0.25 (±0.2)	<0.001	151	0.17 (±0.1)	150	0.23 (±0.2)	<0.001	141	0.16 (±0.2)	139	0.22 (±0.2)	<0.001	52	0.16 (±0.1)	51	0.21 (±0.16)	<0.001
Serum CRP (mg/L)	425	0.17 (±0.6)	422	2.16 (±4.0)	<0.001	275	0.19 (±0.7)	273	2.27 (±4.1)	<0.001	150	0.14 (±0.2)	149	1.96 (±3.8)	<0.001	140	0.14 (±0.2)	138	1.99 (±3.9)	<0.001	51	0.10 (±0.07)	51	1.14 (±1.8)	0.0002
>5mg/L	2	0.5%	54	12.8%	<0.001	2	0.7%	37	13.6%	<0.001	0	0.0%	17	11.4%	N/A	0	0.0%	16	11.6%	N/A	0	0.0%	4	7.8%	N/A
>3mg/L	3	0.7%	83	19.7%	<0.001	3	1.0%	59	21.6%	<0.001	0	0.0%	24	16.1%	N/A	0	0.0%	23	16.7%	N/A	0	0.0%	6	11.8%	N/A
Serum hepcidin (ng/ml)	425	19.9 (±14.2)	417	38.4 (±23.7)	<0.001	277	19.4 (±14.4)	270	38.9 (±23.9)	<0.001	148	20.9 (±13.8)	147	37.4 (±23.5)	<0.001	138	21.5 (±13.9)	136	37.7 (±23.3)	<0.001	52	17.6 (±13.0)	51	33.7 (±22.2)	<0.001
Haemoglobin (g/dl)	414	15.3 (±2.5)	423	19.4 (±3.1)	<0.001	270	15.1 (±2.3)	272	19.1 (±2.9)	<0.001	144	15.6 (±2.7)	151	19.9 (±3.3)	<0.001	136	15.6 (±2.6)	140	19.8 (±3.3)	<0.001	48	16.3 (±3.1)	52	20.7 (±3.5)	<0.001

**Table 3 - Factors associated with neonatal serum iron and TSAT at 6-24h post-partum**

Predictor	Coefficient	Standard error	Standardised coefficient	P value	R2 (%)
Univariate Analysis - Serum Iron					
CRP <sup>a</sup>	0.135	0.023	0.258	<0.0001	15.5
Hepcidin <sup>b</sup>	-0.139	0.025	-0.249	<0.0001	14.2
Transferrin	0.273	0.056	0.222	<0.0001	13.7
Haptoglobin	-1.21	0.303	-0.184	<0.0001	12.2
Age of mother	-0.009	0.002	-0.152	0.001	11.1
Birthweight	-0.0001	-0.00004	-0.149	0.001	11
Multivariate Analysis - Serum Iron					
CRP <sup>a</sup>	0.124	0.022	0.239	<0.0001	29.2
Hepcidin <sup>b</sup>	-0.11	0.025	-0.198	<0.0001	
Transferrin	0.23	0.055	0.186	<0.0001	
Age of mother	-0.008	0.002	-0.143	0.001	
Haptoglobin	-0.741	0.28	-0.114	0.008	
Birthweight	-0.00009	0.00003	-0.104	0.016	
Univariate Analysis - Transferrin Saturation (TSAT)					
CRP <sup>a</sup>	0.153	0.023	0.306	<0.0001	11.7
Haptoglobin	-1.14	0.302	-0.181	<0.0001	5.7
Hepcidin <sup>b</sup>	-0.085	0.025	-0.16	0.001	4.8
Birthweight	-0.0001	0.00004	-0.153	0.001	4.7
Age of mother	-0.008	0.002	-0.151	0.002	4.7
Multivariate Analysis - Transferrin Saturation (TSAT)					
CRP <sup>a</sup>	0.139	0.022	0.285	<0.0001	21.8
Hepcidin <sup>b</sup>	-0.093	0.024	-0.253	<0.0001	
Ferritin	-0.008	0.002	0.187	<0.0001	
Age of mother	-0.68	0.285	-0.161	<0.0001	

**Table 4 - Factors associated with neonatal serum hepcidin at 6-24h post-partum**

Predictor	Coefficient	Standard error	Standardised coefficient	P value	R <sup>2</sup> (%)
<b>Univariate Analysis - Serum Hepcidin</b>					
<b>Ferritin <sup>a</sup></b>	0.039	0.0005	0.354	<0.0001	14
<b>Time of bleed</b>	-0.0007	0.0001	-0.321	<0.0001	12
<b>Transferrin</b>	-0.706	0.104	-0.318	<0.0001	11.5
<b>Serum iron <sup>b</sup></b>	-0.479	0.089	-0.267	<0.0001	8.2
<b>sTfR <sup>c</sup></b>	-0.36	0.092	-0.189	0.001	5
<b>Multivariate Analysis - Serum Hepcidin</b>					
<b>Time of bleed</b>	-0.0008	0.00009	-0.354	<0.0001	36.1
<b>Ferritin <sup>a</sup></b>	0.035	0.004	0.319	<0.0001	
<b>Serum iron <sup>b</sup></b>	-0.413	0.078	-0.232	<0.0001	
<b>sTfR <sup>c</sup></b>	-0.286	0.086	-0.15	0.001	
<b>Transferrin</b>	-0.218	0.111	-0.1	0.05	

**Title**

**Early postnatal hypoferrremia in low birthweight and preterm babies**

**Authors**

James H. Cross, BSc<sup>1</sup>, Ousman Jarjou, MBChB<sup>1</sup>, Nuredin Ibrahim Mohammed, PhD<sup>1</sup>, Santiago Rayment Gomez<sup>2</sup>, Bubacarr J.B Touray, BSc<sup>1</sup>, Andrew M. Prentice, PhD<sup>1</sup>, Carla Cerami, MD<sup>1\*</sup>.

**Affiliations**

<sup>1</sup>Medical Research Council Unit The Gambia at the London School of Hygiene & Tropical Medicine, Atlantic Boulevard, Fajara, P.O. Box 273, Banjul, The Gambia.

<sup>2</sup>Cardiff University, Cardiff, Wales, CF10 3AT, United Kingdom.

## **5.7 SUPPLEMENTARY MATERIAL**

**SUPPLEMENTARY METHODS**

The full study protocol has been previously published.<sup>1</sup>

## SUPPLEMENTARY RESULTS

**Supplementary Table 1 – Participant Characteristics (Groups A-D)**

	Whole Population	Group A (PTB+LBW)	Group B (PTB+NBW)	Group C (FTB+LBW)	Group D (FTB+NBW)	<i>P value</i>
Number of Participants (n)	430	42	99	11	278	
Characteristic	Mean (± SD)					
Gestational age (Weeks)	38.1 (±2.1)	35.1 (±0.9)	35.7 (±0.5)	38.6 (±1.3)	39.4 (±1.3)	<b>&lt;0.0001</b>
Birth weight (g)	3092 (±453.9)	2326 (±122.6)	2916 (±225.9)	2380 (±89.2)	3299 (±368.3)	<b>&lt;0.0001</b>
Head circumference (cm)	34.0 (±1.2)	32.2 (±0.7)	33.4 (±0.8)	32.7 (±1.0)	34.6 (±1.0)	<b>&lt;0.0001</b>
Length (cm)	49.2 (±1.9)	46.6 (±1.4)	48.2 (±1.4)	47.0 (±1.5)	50.0 (±1.6)	<b>&lt;0.0001</b>
Maternal hemoglobin ≤7 days before delivery (g/dl)	11.6 (±1.8)	12.2 (±1.5)	11.3 (±1.8)	10.8 (±2.0)	11.6 (±1.8)	0.29
Age of mother (Years)	29.4 (±6.9)	26.9 (±6.6)	29.8 (±6.7)	26.4 (±6.9)	29.7 (±6.9)	<b>0.03</b>
1 min APGAR score (0-10)	9.65 (±0.7)	9.6 (±0.7)	9.8 (±0.48)	9.2 (±1.3)	9.6 (±0.8)	<b>0.03</b>
Time from admission to birth (Hours)	3.23 (±5.1)	3.2 (±4.9)	3.2 (±4.8)	3.7 (±5.5)	3.2 (±5.2)	0.99
Time from delivery to V1 blood collection (Hours)	12.7 (±5.3)	13.0 (±4.9)	13.0 (±5.5)	10.5 (±3.8)	12.7 (±5.4)	0.52
	Percentage (%)					
Percentage male (%)	53.5% (224)	47.6% (20)	52.5% (52)	63.6% (7)	54.3% (151)	0.76
Percentage early term, ≥37-≤38 weeks (%)	19.3% (83)	0% (0)	0% (0)	54.5% (6)	27.7% (77)	<b>&lt;0.0001</b>
G6PD deficiency positive (%)	12.6% (54)	7.1% (3)	18.2% (18)	9.1% (1)	11.5% (32)	0.23
Multiple births (%)	3% (13)	7.1% (3)	4% (4)	0% (0)	2.2% (6)	0.28
Percentage of mother on antenatal iron/folic acid (%)	82.1% (353)	76.2% (32)	87.9% (87)	63.6% (7)	81.7% (227)	0.21

**Supplementary Table 1:** Baseline Characteristics. Demographic, clinical and pregnancy outcome characteristics of the women and their newborns for groups A-D. Data are presented as mean (± SD) and analysed by ANOVA or as n (%) and analysed by  $\chi^2$  test. *P* values in bold font are considered significant based on  $P < 0.05$ .

**Supplementary Table 2 – Maternal Ethnicity**

Ethnic Group	Whole Population	FTB/NBW	PTB/LBW	<i>P value</i>	Full-Term Birth (FTB)	Preterm Birth (PTB)	<i>P value</i>	NBW	LBW	<i>P value</i>	Group A (PTB+LBW)	Group B (PTB+NBW)	Group C (FTB+LBW)	Group D (FTB+NBW)	<i>P value</i>
<b>Aku</b>	0.9% (4)	1.4% (4)	0% (0)		1.4% (4)	0% (0)		1.1% (4)	0% (0)		0% (0)	0% (0)	0% (0)	1.4% (4)	
<b>Balanta</b>	0.9% (4)	0.4% (4)	2% (3)		0.4% (1)	2.1% (3)		0.8% (3)	1.9% (1)		2.4% (1)	2% (2)	0% (0)	0.4% (1)	
<b>Fula</b>	18.4% (79)	17.3% (79)	20.4% (31)		17% (49)	21.3% (30)		18.6% (70)	17% (9)		19.1% (8)	22.2% (22)	9.1% (1)	17.3% (48)	
<b>Jola</b>	13.3% (57)	14.8% (41)	10.5% (16)		14.5% (42)	10.6% (15)		13.5% (51)	11.3% (6)		11.9% (5)	10.1% (10)	9.1% (1)	14.8% (41)	
<b>Mandinka</b>	38.4% (165)	38.9% (108)	37.5% (57)		39.5% (114)	36.2% (51)		38.7% (146)	35.9% (19)		31% (13)	38.4% (38)	54.6% (6)	38.9% (108)	
<b>Manjago</b>	0.7% (3)	1.1% (3)	0% (0)	0.13	1% (3)	0% (0)	0.1	0.8% (3)	0% (0)	0.73	0% (0)	0% (0)	0% (0)	1.1% (3)	0.65
<b>Sarahule</b>	5.1% (22)	3.6% (10)	7.9% (12)		3.8% (11)	7.8% (11)		4.5% (17)	9.4% (5)		9.5% (4)	7% (7)	9.1% (1)	3.6% (10)	
<b>Serere</b>	6.3% (27)	5.4% (15)	7.9% (12)		5.2% (15)	8.5% (12)		6.6% (25)	3.8% (2)		4.8% (2)	10.1% (10)	0% (0)	5.4% (15)	
<b>Wollof</b>	11.9% (51)	13% (36)	9.9% (15)		13.2% (38)	9.2% (13)		11.1% (42)	17% (9)		16.7% (7)	6% (6)	18.2% (2)	13% (36)	
<b>Other</b>	4.2% (18)	4.3% (12)	4% (6)		4.2% (12)	4.3% (6)		4.2% (16)	3.8% (2)		4.8% (2)	4% (4)	0% (0)	4.3% (12)	

**Supplementary Table 2:** Ethnicity of mother. Data are presented as n (%) and analysed by  $\chi^2$  test. *P* values in bold font are considered significant based on  $P < 0.05$ . No significant differences between groups was observed.

**Supplementary Table 3 – Comparison of iron status, inflammation and haematological parameters in cord samples**

	Whole Population		PTB/LBW		FTB/NBW			Full-Term Birth (FTB)		Preterm Birth (PTB)			Normal Birthweight (NBW)		Low Birthweight (LBW)		
Sample Type	n	Cord	n	Cord	n	Cord	P value	n	Cord	n	Cord	P value	n	Cord	n	Cord	P value
Serum iron (µmol/L)	425	23.3 (±7.1)	150	24.3 (±7.3)	275	22.7 (±7.0)	<b>0.02</b>	285	22.7 (±7.0)	140	24.4 (±7.3)	<b>0.01</b>	374	23.1 (±7.0)	51	24.5 (±7.7)	0.2
UIBC (µmol/L)	423	22.9 (±10.5)	150	21.5 (±10.6)	273	23.7 (±10.4)	<b>0.04</b>	283	23.8 (±10.8)	140	21.2 (±9.8)	<b>0.02</b>	372	23.0 (±10.3)	51	22.1 (±12.2)	0.6
TIBC (µmol/L)	423	46.1 (±8.0)	150	45.8 (±7.8)	273	46.3 (±8.1)	0.3	283	46.4 (±8.3)	140	45.6 (±7.4)	0.3	372	46.1 (±7.8)	51	46.6 (±9.3)	0.6
Transferrin (g/L)	426	1.96 (±0.3)	151	1.93 (±0.3)	275	1.98 (±0.3)	0.1	285	1.99 (±0.4)	141	1.92 (±0.3)	<b>0.05</b>	374	1.96 (±0.3)	52	1.94 (±0.4)	0.6
TSAT (%)	423	51.7 (±17.3)	150	54.5 (±18.0)	273	50.2 (±16.7)	<b>0.01</b>	283	50.2 (±16.9)	140	54.8 (±17.7)	<b>0.01</b>	372	51.4 (±17.0)	51	54.6 (±18.7)	0.2
Serum ferritin (ug/L)	426	210.5 (±157.8)	151	206.6 (±158.5)	275	212.6 (±157.7)	0.7	285	212.3 (±159.6)	141	206.8 (±154.8)	0.7	374	213.0 (±154.7)	52	192.1 (±179.5)	0.4
Serum AGP (g/L)	426	0.18 (±0.1)	151	0.17 (±0.2)	275	0.19 (±0.1)	0.2	285	0.19 (±0.1)	141	0.16 (±0.2)	0.08	374	0.18 (±0.1)	52	0.16 (±0.1)	0.3
Serum CRP (mg/L)	425	0.17 (±0.6)	150	0.14 (±0.3)	275	0.19 (±0.7)	0.4	285	0.19 (±0.7)	140	0.14 (±0.3)	0.4	374	0.18 (±0.6)	51	0.10 (±0.07)	0.3
>5mg/L	2	0.5%	0	0	2	0.7%	N/A	2	0.7%	0	0	N/A	2	0.5%	0	0	N/A
>3mg/L	3	0.7%	0	0	3	1.1%	N/A	3	1.1%	0	0	N/A	3	0.8%	0	0	N/A
Serum hepcidin (ng/ml)	425	19.9 (±14.2)	148	20.9 (±13.8)	277	19.4 (±14.4)	0.3	287	19.1 (±14.3)	138	21.5 (±13.9)	0.1	373	20.2 (±14.3)	52	17.6 (±13.0)	0.2
Haemoglobin (g/dl)	414	15.3 (±2.5)	144	15.6 (±2.7)	270	15.1 (±2.3)	<b>0.04</b>	278	15.1 (±2.4)	136	15.6 (±2.6)	0.06	366	15.1 (±2.4)	48	16.3 (±3.1)	<b>0.003</b>
Haematocrit (%)	414	42.7 (±7.4)	144	43.8 (±8.1)	270	42.2 (±7.0)	<b>0.039</b>	278	42.3 (±7.2)	136	43.7 (±7.8)	0.06	366	42.3 (±7.1)	48	45.9 (±9.2)	<b>0.0016</b>
Soluble transferrin receptor (mg/L)	424	6.0 (±2.0)	151	5.94 (±1.9)	273	5.99 (±2.0)	0.79	283	6.0 (±2.1)	141	5.9 (±1.7)	0.41	372	5.92 (±1.9)	52	6.3 (±2.4)	0.19
Haptoglobin (g/L)	426	0.02 (±0.07)	151	0.03 (±0.09)	275	0.02 (±0.06)	0.31	285	0.02 (±0.06)	141	0.03 (±0.09)	0.28	374	0.02 (±0.08)	52	0.01 (±0.02)	0.28
Undetectable haptoglobin (%)	182	42.7%	66	43.8%	116	42.2%	0.76	120	42.1%	63	44.6%	0.71	158	42.2%	24	46.2%	0.307
Mean corpuscular volume (fl)	414	98.0 (±6.2)	144	98.5 (±6.1)	270	97.8 (±6.3)	0.23	278	97.7 (±6.4)	136	98.7 (±5.7)	0.11	366	97.9 (±6.0)	48	99.3 (±7.3)	0.15
Mean corpuscular haemoglobin (pg)	414	35.1 (±2.4)	144	35.3 (±2.5)	270	35.0 (±2.4)	0.38	278	35.0 (±2.5)	136	35.4 (±2.3)	0.19	366	35.1 (±2.4)	48	35.3 (±3.0)	0.61
Mean corpuscular haemoglobin concentration (g/dl)	414	35.8 (±1.0)	144	35.8 (±1.0)	270	35.9 (±1.0)	0.5	278	35.8 (±1.0)	136	35.8 (±1.0)	0.68	366	35.9 (±0.9)	48	35.5 (±0.9)	<b>0.027</b>
White blood cell count (unit/L)	412	13.0 (±5.3)	143	12.3 (±4.6)	269	13.3 (±5.7)	0.07	277	13.3 (±5.7)	135	12.2 (±4.4)	<b>0.033</b>	364	13.2 (±5.4)	48	11.5 (±4.8)	<b>0.048</b>
Lymphocyte count (unit/L)	412	4.73 (±3.1)	143	4.67 (±2.2)	269	4.77 (±3.5)	0.74	277	4.82 (±3.5)	135	4.54 (±2.0)	0.39	364	4.72 (±3.1)	48	4.84 (±2.4)	0.79
Lymphocyte percentage (%)	412	36.2 (±8.3)	143	38.1 (±8.7)	269	35.2 (±7.9)	<b>0.0006</b>	277	35.4 (±8.0)	135	37.8 (±8.7)	<b>0.0063</b>	364	35.5 (±7.9)	48	42.0 (±8.8)	<b>&lt;0.001</b>
MID cell count (unit/L)	412	1.02 (±0.6)	143	0.99 (±0.5)	269	1.05 (±0.6)	0.35	277	1.05 (±0.6)	135	0.97 (±0.5)	0.22	364	1.0 (±0.6)	48	1.04 (±0.6)	0.82
MID cell percentage (%)	412	8.27 (±2.6)	143	8.52 (±3.0)	269	8.15 (±2.4)	0.17	277	8.17% (±2.5)	135	8.48 (±3.0)	0.25	364	8.12 (±2.3)	48	9.46 (±4.2)	<b>0.0008</b>



Granulocyte count (unit/L)	412	7.21 (±2.7)	143	6.68 (±2.8)	269	7.49 (±2.7)	0.0039	277	7.47 (±2.7)	135	6.66 (±2.8)	0.0047	364	7.41 (±2.7)	48	5.67 (±2.7)	<0.001	
Granulocyte percentage (%)	412	55.5 (±9.1)	143	53.3 (±9.4)	269	56.6 (±8.7)	0.0004	277	56.4 (±8.8)	135	53.7 (±9.4)	0.0045	364	56.4 (±8.6)	48	48.5 (±9.8)	<0.001	
Red blood cell count (unit/L)	414	4.36 (±0.7)	144	4.45 (±0.8)	270	4.32 (±0.7)	0.096	278	4.33 (±0.7)	136	4.42 (±0.8)	0.2	366	4.33 (±0.7)	48	4.63 (±0.9)	0.0066	
Red blood cell distribution width (%)	414	15.2 (±1.1)	144	15.2 (±1.0)	270	15.2 (±1.2)	0.97	278	15.2 (±1.2)	136	15.1 (±1.0)	0.64	366	15.2 (±1.1)	48	15.2 (±0.9)	0.95	
Red blood cell distribution width - absolute (fl)	414	81.0 (±8.6)	144	81.5 (±8.1)	270	80.7 (±8.9)	0.39	278	80.7 (±9.0)	136	81.6 (±7.8)	0.32	366	80.8 (±8.6)	48	82.0 (±8.7)	0.39	
Platelet count (unit/L)	414	241.4 (±86.9)	144	236.7 (±74.8)	270	243.8 (±92.8)	0.43	278	243.2 (±93.4)	136	237.5 (±72.1)	0.53	366	243.3 (±87.3)	48	225.8 (±83.1)	0.19	
Mean platelet volume (fl)	409	8.34 (±0.8)	142	8.30 (±0.8)	267	8.37 (±0.8)	0.38	275	8.36 (±0.8)	134	8.3 (±0.8)	0.45	362	8.36 (±0.8)	47	8.17 (±0.8)	0.094	
Platelet distribution width (%)	409	43.3 (±2.8)	142	43.0 (±2.5)	267	43.5 (±3.0)	0.063	275	43.4 (±3.0)	134	42.9 (±2.5)	0.091	362	43.3 (±2.8)	47	42.9 (±2.8)	0.39	
Platelet distribution width - absolute (fl)	409	11.8 (±1.2)	142	11.7 (±1.1)	267	11.8 (±1.3)	0.17	275	11.8 (±1.3)	134	11.7 (±1.1)	0.22	362	11.8 (±1.2)	47	11.5 (±1.1)	0.11	
Platelet crit (%) <sup>a</sup>	409	0.2 (±0.06)	142	0.194 (±0.06)	267	0.201 (±0.07)	0.24	275	0.2 (±0.06)	134	0.194 (±0.05)	0.36	362	0.201 (±0.06)	47	0.18 (±0.06)	0.057	
Plate large cell ratio (%)	409	17.8 (±5.3)	142	17.5 (±5.1)	267	18.0 (±5.4)	0.3	275	18.0 (±5.3)	134	17.5 (±5.1)	0.31	362	18.0 (±5.3)	47	16.8 (±4.9)	0.14	
Plate large cell count (unit/L)	409	41.4 (±14.2)	142	40.1 (±13.2)	267	42.1 (±14.6)	0.16	275	41.9 (±14.8)	134	40.2 (±12.7)	0.24	362	42.0 (±14.1)	47	36.6 (±13.7)	0.014	
Haemolysis score (0-6)																		
	0	166	39.0%	54	35.8%	112	41.2%	0.27	116	41.1%	50	35.5%	0.26	149	40.2%	17	32.7%	0.3
	1	144	34.0%	53	35.1%	91	33.5%	0.73	93	33.0%	51	36.2%	0.51	132	35.6%	12	23.1%	0.074
	2	78	18.4%	29	19.2%	49	18.00%	0.6	52	18.4%	26	18.4%	0.99	62	16.7%	16	30.8%	0.014
	3	35	8.3%	15	9.93%	20	7.35%	0.4	21	7.5%	14	9.9%	0.38	28	7.6%	7	13.5%	0.15
	4	0	0.0%	0	0.00%	0	0.00%	..	0	0.0%	0	0.0%	..	0	0.00%	0	0.00%	..

**Supplementary Table 3:** Further Iron, Infection and Haematological Parameters (umbilical cord samples) for FTB/NBW, PTB/LBW, FTB, PTB, NBW and LBW neonates. Data are presented as mean (± SD) and analysed by ANOVA. *P* values in bold font are considered significant based on *P*<0.05. <sup>a</sup> = Platelet Crit (%) is a measure of total platelet mass.

**Supplementary Table 4 – Comparison of iron status, inflammation and haematological parameters in venous samples**

	Whole Population		PTB/LBW		FTB/NBW			Full-Term Birth (FTB)		Preterm Birth (PTB)			Normal Birthweight (NBW)		Low Birthweight (LBW)		
Sample Type	n	Venous	n	Venous	n	Venous	P value	n	Venous	n	Venous	P value	n	Venous	n	Venous	P value
Serum iron (µmol/L)	421	7.5 (±4.5)	150	8.0 (±4.1)	271	7.3 (±4.6)	0.07	282	7.4 (±4.6)	139	7.8 (±4.1)	0.2	370	7.4 (±4.6)	51	8.5 (±3.4)	<b>0.05</b>
UIBC (µmol/L)	420	43.5 (±15.9)	149	42.2 (±10.9)	271	44.1 (±18.0)	0.2	282	44.2 (±17.8)	138	42.0 (±10.9)	0.2	370	43.4 (±16.5)	50	43.8 (±10.2)	0.9
TIBC (µmol/L)	420	51.0 (±17.9)	149	50.2 (±11.3)	271	51.4 (±20.7)	0.5	282	51.5 (±20.5)	138	49.8 (±11.1)	0.4	370	50.8 (±18.6)	50	52.3 (±11.9)	0.2
Transferrin (g/L)	423	1.99 (±0.3)	150	2.00 (±0.3)	273	1.98 (±0.3)	0.6	284	1.99 (±0.3)	139	1.99 (±0.3)	0.8	372	1.99 (±0.3)	51	2.02 (±0.4)	0.5
TSAT (%)	420	15.0 (±6.9)	149	16.2 (±8.2)	271	14.4 (±6.1)	<b>0.01</b>	282	14.5 (±6.0)	138	16.1 (±8.4)	<b>0.01</b>	370	14.9 (±7.1)	50	16.2 (±5.3)	0.2
Serum ferritin (ug/L)	415	381.2 (±297.6)	149	358.6 (±268.1)	271	393.9 (±312.6)	0.3	277	393.9 (±314.5)	138	355.7 (±259.6)	0.2	364	383.3 (±291.0)	51	366.2 (±343.8)	0.7
Serum AGP (g/L)	423	0.24 (±0.2)	150	0.23 (±0.2)	273	0.25 (±0.2)	0.2	284	0.25 (±0.17)	139	0.22 (±0.2)	0.2	372	0.25 (±0.2)	51	0.21 (±0.2)	0.1
Serum CRP (mg/L)	422	2.16 (±4.0)	149	1.96 (±3.8)	273	2.27 (±4.1)	0.5	284	2.24 (±4.1)	138	1.99 (±3.9)	0.5	371	2.3 (±4.2)	51	1.14 (±1.8)	<b>0.05</b>
>5mg/L	54	12.8%	17	11.4%	37	13.6%	0.5	38	13.4%	16	11.6%	0.6	50	13.5%	4	7.8%	0.3
>3mg/L	83	19.7%	24	16.1%	59	21.6%	0.2	60	21.1%	23	16.7%	0.3	77	20.8%	6	11.8%	0.1
Serum hepcidin (ng/ml)	417	38.4 (±23.7)	147	37.4 (±23.5)	270	38.9 (±23.9)	0.5	281	38.7 (±24.0)	136	37.7 (±23.3)	0.7	366	39.0 (±23.9)	51	33.7 (±22.2)	0.1
Haemoglobin (g/dl)	423	19.4 (±3.1)	151	19.9 (±3.3)	272	19.1 (±2.9)	<b>0.02</b>	283	19.2 (±3.0)	140	19.8 (±3.3)	<b>0.04</b>	371	19.2 (±3.0)	52	20.7 (±3.5)	<b>0.001</b>
Haematocrit (%)	423	54.7 (±9.3)	151	56.1 (±10.1)	272	53.9 (±8.9)	<b>0.025</b>	283	54.1 (±9.0)	140	55.9 (±9.8)	0.64	371	54.1 (±9.0)	52	59.0 (±10.7)	<b>0.0004</b>
Soluble transferrin receptor (mg/L)	421	7.0 (±2.2)	150	7.03 (±2.1)	271	6.94 (±2.2)	0.7	282	7.0 (±2.3)	139	6.9 (±2.0)	0.64	370	6.87 (±2.1)	51	7.71 (±2.5)	<b>0.0093</b>
Haptoglobin (g/L)	423	0.02 (±0.06)	150	0.02 (±0.04)	273	0.03 (±0.07)	0.35	284	0.03 (±0.07)	139	0.02 (±0.04)	0.41	372	0.02 (±0.07)	51	0.01 (±0.01)	0.25
Undetectable haptoglobin (%)	144	34.0%	43	28.7%	101	37.00%	0.083	106	37.3%	38	27.3%	<b>0.041</b>	129	34.7%	15	29.4%	0.46
Mean corpuscular volume (fl)	423	98.0 (±6.1)	151	98.5 (±6.2)	273	97.7 (±6.0)	0.21	283	97.6 (±6.2)	140	98.8 (±5.8)	<b>0.048</b>	371	97.9 (±5.8)	52	98.9 (±7.6)	0.26
Mean corpuscular haemoglobin (pg)	423	34.9 (±2.4)	151	35.0 (±2.4)	272	34.8 (±2.4)	0.48	283	34.8 (±2.5)	140	35.1 (±2.3)	0.13	371	34.9 (±2.4)	52	34.8 (±2.9)	0.82
Mean corpuscular haemoglobin concentration (g/dl)	423	35.6 (±1.0)	151	35.5 (±0.9)	272	35.6 (±1.1)	0.39	283	35.6 (±1.1)	140	35.5 (±0.9)	0.59	371	35.6 (±1.0)	52	35.2 (±0.9)	<b>0.0036</b>
White blood cell count (unit/L)	422	15.9 (±5.5)	151	14.9 (±4.7)	271	16.4 (±5.8)	<b>0.0068</b>	282	16.3 (±5.7)	140	15.0 (±4.8)	<b>0.019</b>	370	16.3 (±5.5)	52	13.3 (±4.1)	<b>0.0002</b>
Lymphocyte count (unit/L)	422	4.1 (±2.4)	151	3.93 (±1.4)	271	4.21 (±2.9)	0.25	282	4.19 (±2.8)	140	3.95 (±1.4)	0.35	370	4.17 (±2.6)	52	3.72 (±1.2)	0.22
Lymphocyte percentage (%)	422	26.9 (±8.8)	151	27.8 (±8.9)	271	26.5 (±8.8)	0.14	282	26.5 (±8.7)	140	27.8 (±9.0)	0.14	370	26.5 (±8.7)	52	29.7 (±9.2)	<b>0.014</b>
MID cell count (unit/L)	422	1.48 (±1.0)	151	1.37 (±0.9)	271	1.55 (±1.1)	0.08	282	1.54 (±1.0)	140	1.37 (±0.9)	0.11	370	1.52 (±1.0)	52	1.27 (±0.9)	0.097
MID cell percentage (%)	422	9.67 (±5.4)	151	9.43 (±5.0)	271	9.81 (±5.6)	0.49	282	9.79 (±5.6)	140	9.41 (±5.0)	0.49	370	9.67 (±5.4)	52	9.75 (±5.1)	0.91

Granulocyte count (unit/L)	422	10.3 (±4.1)	151	9.63 (±3.8)	271	10.67 (±4.2)	0.012	282	10.6 (±4.1)	140	9.68 (±3.9)	0.029	370	10.57 (±4.1)	52	8.31 (±3.3)	0.0002	
Granulocyte percentage (%)	422	63.4 (±8.6)	151	62.8 (±8.1)	271	63.7 (±8.8)	0.28	282	63.7 (±8.7)	140	62.7 (±8.2)	0.27	370	63.8 (±8.5)	52	60.5 (±8.2)	0.0095	
Red blood cell count (unit/L)	423	5.58 (±1.0)	151	5.69 (±1.0)	272	5.53 (±0.9)	0.093	283	5.55 (±1.0)	140	5.66 (±0.9)	0.28	371	5.54 (±0.9)	52	5.97 (±1.1)	0.0018	
Red blood cell distribution width (%)	423	15.4 (±1.3)	151	15.4 (±1.0)	272	15.4 (±1.4)	0.49	283	15.5 (±1.4)	140	15.3 (±1.0)	0.36	371	15.4 (±1.3)	52	15.3 (±0.9)	0.6	
Red blood cell distribution width - Absolute (fl)	423	82.2 (±8.5)	151	82.6 (±8.4)	272	82.0 (±8.6)	0.47	283	81.9 (±8.8)	140	82.8 (±8.1)	0.26	371	82.1 (8.4)	52	82.8 (±9.2)	0.58	
Platelet count (unit/L)	423	260.4 (±94.6)	151	263.4 (±96.0)	272	258.7 (±94.0)	0.62	283	257.8 (±93.6)	140	265.7 (±96.7)	0.42	371	261.6 (±95.9)	52	252.1 (±85.4)	0.5	
Mean platelet volume (fl)	419	8.47 (±0.8)	148	8.38 (±0.7)	272	8.52 (±0.8)	0.064	282	8.52 (±0.8)	137	8.37 (±0.7)	0.061	367	8.49 (±0.8)	52	8.37 (±0.8)	0.31	
Platelet distribution width (%)	419	43.7 (±3.2)	148	43.4 (±2.8)	271	43.9 (±3.3)	0.082	282	44.0 (±3.3)	137	43.3 (±2.8)	0.051	367	43.8 (±3.2)	52	42.4 (±3.0)	0.39	
Platelet distribution width - Absolute (fl)	419	12.0 (±1.3)	148	11.8 (±1.1)	271	12.1 (±1.4)	0.045	282	12.1 (±1.4)	137	11.8 (±1.1)	0.035	367	12.0 (±1.3)	52	11.8 (±1.3)	0.22	
Platelet crit (%)	419	0.22 (±0.07)	148	0.22 (±0.07)	271	0.215 (±0.07)	0.59	282	0.214 (±0.07)	137	0.221 (±0.07)	0.33	367	0.218 (±0.07)	52	0.207 (±0.06)	0.32	
Plate large cell ratio (%)	419	18.8 (±5.5)	148	18.1 (±4.9)	271	19.2 (±5.8)	0.045	282	19.2 (±5.8)	137	18.0 (±4.9)	0.032	367	18.9 (±5.6)	52	18.2 (±5.3)	0.38	
Plate large cell count (unit/L)	419	46.5 (±15.8)	148	45.9 (±13.4)	271	46.9 (±17.0)	0.55	282	46.7 (±16.9)	137	46.3 (±13.4)	0.79	367	47.0 (±16.3)	52	43.6 (±11.6)	0.15	
Haemolysis score (0-6)																		
	0	0	0.0%	17	32.7%	33	12.2%	<0.001	0	0.0%	0	0.0%	..	39	10.5%	4	7.8%	0.55
	1	43	10.2%	12	23.1%	135	49.9%	<0.001	34	12.1%	9	6.5%	0.075	192	51.9%	21	41.2%	0.15
	2	213	50.6%	16	30.7%	92	34.00%	0.66	138	49.0%	75	54.0%	0.33	125	33.8%	25	49.00%	0.033
	3	150	35.6%	7	13.5%	11	4.1%	0.006	99	35.1%	51	36.7%	0.75	14	3.8%	1	1.96%	0.51
	4	15	3.5%	0	0.00%	0	0.00%	..	11	3.9%	4	2.9%	0.59	0	0.00%	0	0.00%	..

**Supplementary Table 4:** Further Iron, Infection and Haematological Parameters (venous samples) for FTB/NBW, PTB/LBW, FTB, PTB, NBW and LBW neonates. Data are presented as mean (± SD) and analysed by ANOVA. *P* values in bold font are considered significant based on *P* < 0.05.

**Supplementary Table 5 – Changes in further iron status and inflammation variables between cord and post-natal blood**

	Whole Population					FTB/NBW					PTB/LBW					Preterm Birth (PTB)					Low Birthweight (LBW)				
Sample Type	n	Cord	n	Venous	P value	n	Cord	n	Venous	P value	n	Cord	n	Venous	P value	n	Cord	n	Venous	P value	n	Cord	n	Venous	P value
Hematocrit (%)	414	42.7 (±7.4)	423	54.7 (±9.3)	<0.001	270	42.2 (±7.0)	272	53.9 (±8.9)	<0.001	144	43.8 (±8.1)	151	56.1 (±10.1)	<0.001	136	43.7 (±7.8)	140	55.9 (±9.8)	<0.001	48	45.9 (±9.2)	52	59.0 (±10.7)	<0.001
Soluble transferrin receptor (mg/L)	424	6.0 (±2.0)	421	7.0 (±2.2)	<0.001	273	5.99 (±2.0)	271	6.94 (±2.2)	<0.001	151	5.94 (±1.9)	150	7.03 (±2.1)	<0.001	141	5.9 (±1.7)	139	6.9 (±2.0)	<0.001	52	6.3 (±2.4)	51	7.71 (±2.5)	<0.001
Haptoglobin (g/L)	426	0.02 (±0.07)	423	0.02 (±0.06)	0.52	275	0.02 (±0.06)	273	0.03 (±0.07)	0.06	151	0.03 (±0.09)	150	0.02 (±0.04)	0.19	141	0.03 (±0.09)	139	0.02 (±0.04)	0.2	52	0.01 (±0.02)	51	0.01 (±0.01)	0.28
Undetectable haptoglobin (%)	182	42.7%	144	34.0%	0.009	116	42.2%	101	37.00%	0.21	66	43.8%	43	28.7%	0.006	63	44.6%	38	27.3%	0.002	24	46.2%	15	29.4%	0.08
Mean corpuscular volume (fl)	414	98.0 (±6.2)	423	98.0 (±6.1)	0.63	270	97.8 (±6.3)	273	97.7 (±6.0)	0.29	144	98.5 (±6.1)	151	98.5 (±6.2)	0.42	136	98.7 (±5.7)	140	98.8 (±5.8)	0.24	48	99.3 (±7.3)	52	98.9 (±7.6)	0.37
Mean corpuscular haemoglobin (pg)	414	35.1 (±2.4)	423	34.9 (±2.4)	<0.001	270	35.0 (±2.4)	272	34.8 (±2.4)	<0.001	144	35.3 (±2.5)	151	35.0 (±2.4)	<0.001	136	35.4 (±2.3)	140	35.1 (±2.3)	<0.001	48	35.3 (±3.0)	52	34.8 (±2.9)	<0.001
Mean corpuscular haemoglobin concentration (g/dl)	414	35.8 (±1.0)	423	35.6 (±1.0)	<0.001	270	35.9 (±1.0)	272	35.6 (±1.1)	<0.001	144	35.8 (±1.0)	151	35.5 (±0.9)	0.0001	136	35.8 (±1.0)	140	35.5 (±0.9)	<0.001	48	35.5 (±0.9)	52	35.2 (±0.9)	0.01
White blood cell count (unit/L)	412	13.0 (±5.3)	422	15.9 (±5.5)	<0.001	269	13.3 (±5.7)	271	16.4 (±5.8)	<0.001	143	12.3 (±4.6)	151	14.9 (±4.7)	<0.001	135	12.2 (±4.4)	140	15.0 (±4.8)	<0.001	48	11.5 (±4.8)	52	13.3 (±4.1)	0.008
Lymphocyte count (unit/L)	412	4.73 (±3.1)	422	4.1 (±2.4)	<0.001	269	4.77 (±3.5)	271	4.21 (±2.9)	0.0001	143	4.67 (±2.2)	151	3.93 (±1.4)	0.0001	135	4.54 (±2.0)	140	3.95 (±1.4)	0.0006	48	4.84 (±2.4)	52	3.72 (±1.2)	0.003
Lymphocyte percentage (%)	412	36.2 (±8.3)	422	26.9 (±8.8)	<0.001	269	35.2 (±7.9)	271	26.5 (±8.8)	<0.001	143	38.1 (±8.7)	151	27.8 (±8.9)	<0.001	135	37.8 (±8.7)	140	27.8 (±9.0)	<0.001	48	42.0 (±8.8)	52	29.7 (±9.2)	<0.001
MID cell count (unit/L)	412	1.02 (±0.6)	422	1.48 (±1.0)	<0.001	269	1.05 (±0.6)	271	1.55 (±1.1)	<0.001	143	0.99 (±0.5)	151	1.37 (±0.9)	<0.001	135	0.97 (±0.5)	140	1.37 (±0.9)	<0.001	48	1.04 (±0.6)	52	1.27 (±0.9)	0.1
MID cell percentage (%)	412	8.27 (±2.6)	422	9.67 (±5.4)	<0.001	269	8.15 (±2.4)	271	9.81 (±5.6)	<0.001	143	8.52 (±3.0)	151	9.43 (±5.0)	0.04	135	8.48 (±3.0)	140	9.41 (±5.0)	0.04	48	9.46 (±4.2)	52	9.75 (±5.1)	0.6
Granulocyte count (unit/L)	412	7.21 (±2.7)	422	10.3 (±4.1)	<0.001	269	7.49 (±2.7)	271	10.67 (±4.2)	<0.001	143	6.68 (±2.8)	151	9.63 (±3.8)	<0.001	135	6.66 (±2.8)	140	9.68 (±3.9)	<0.001	48	5.67 (±2.7)	52	8.31 (±3.3)	<0.001
Granulocyte percentage (%)	412	55.5 (±9.1)	422	63.4 (±8.6)	<0.001	269	56.6 (±8.7)	271	63.7 (±8.8)	<0.001	143	53.3 (±9.4)	151	62.8 (±8.1)	<0.001	135	53.7 (±9.4)	140	62.7 (±8.2)	<0.001	48	48.5 (±9.8)	52	60.5 (±8.2)	<0.001
Red blood cell count (unit/L)	414	4.36 (±0.7)	423	5.58 (±1.0)	<0.001	270	4.32 (±0.7)	272	5.53 (±0.9)	<0.001	144	4.45 (±0.8)	151	5.69 (±1.0)	<0.001	136	4.42 (±0.8)	140	5.66 (±0.9)	<0.001	48	4.63 (±0.9)	52	5.97 (±1.1)	<0.001
Red blood cell distribution width (%)	414	15.2 (±1.1)	423	15.4 (±1.3)	<0.001	270	15.2 (±1.2)	272	15.4 (±1.4)	<0.001	144	15.2 (±1.0)	151	15.4 (±1.0)	<0.001	136	15.1 (±1.0)	140	15.3 (±1.0)	<0.001	48	15.2 (±0.9)	52	15.3 (±0.9)	0.002
Red blood cell distribution width - absolute (fl)	414	81.0 (±8.6)	423	82.2 (±8.5)	<0.001	270	80.7 (±8.9)	272	82.0 (±8.6)	0.0001	144	81.5 (±8.1)	151	82.6 (±8.4)	<0.001	136	81.6 (±7.8)	140	82.8 (±8.1)	<0.001	48	82.0 (±8.7)	52	82.8 (±9.2)	<0.001
Platelet count (unit/L)	414	241.4 (±86.9)	423	260.4 (±94.6)	0.0003	270	243.8 (±92.8)	272	258.7 (±94.0)	0.022	144	236.7 (±74.8)	151	263.4 (±96.0)	0.003	136	237.5 (±72.1)	140	265.7 (±96.7)	0.003	48	225.8 (±83.1)	52	252.1 (±85.4)	0.09
Mean platelet volume (fl)	409	8.34 (±0.8)	419	8.47 (±0.8)	0.0003	267	8.37 (±0.8)	272	8.52 (±0.8)	0.0001	142	8.30 (±0.8)	148	8.38 (±0.7)	0.37	134	8.3 (±0.8)	137	8.37 (±0.7)	0.29	47	8.17 (±0.8)	52	8.37 (±0.8)	0.07
Platelet distribution width (%)	409	43.3 (±2.8)	419	43.7 (±3.2)	0.007	267	43.5 (±3.0)	271	43.9 (±3.3)	0.021	142	43.0 (±2.5)	148	43.4 (±2.8)	0.16	134	42.9 (±2.5)	137	43.3 (±2.8)	0.26	47	42.9 (±2.8)	52	42.4 (±3.0)	0.61
Platelet distribution width - absolute (fl)	409	11.8 (±1.2)	419	12.0 (±1.3)	0.0006	267	11.8 (±1.3)	271	12.1 (±1.4)	0.0009	142	11.7 (±1.1)	148	11.8 (±1.1)	0.24	134	11.7 (±1.1)	137	11.8 (±1.1)	0.3	47	11.5 (±1.1)	52	11.8 (±1.3)	0.21
Platelet crit (%)	409	0.2 (±0.06)	419	0.22 (±0.07)	<0.001	267	0.201 (±0.07)	271	0.215 (±0.07)	0.01	142	0.194 (±0.06)	148	0.22 (±0.07)	0.0001	134	0.194 (±0.05)	137	0.221 (±0.07)	0.0001	47	0.18 (±0.06)	52	0.207 (±0.06)	0.034
Plate large cell ratio (%)	409	17.8 (±5.3)	419	18.8 (±5.5)	0.0001	267	18.0 (±5.4)	271	19.2 (±5.8)	0.0001	142	17.5 (±5.1)	148	18.1 (±4.9)	0.27	134	17.5 (±5.1)	137	18.0 (±4.9)	0.32	47	16.8 (±4.9)	52	18.2 (±5.3)	0.06
Plate large cell count (unit/L)	409	41.4 (±14.2)	419	46.5 (±15.8)	<0.001	267	42.1 (±14.6)	271	46.9 (±17.0)	<0.001	142	40.1 (±13.2)	148	45.9 (±13.4)	<0.001	134	40.2 (±12.7)	137	46.3 (±13.4)	<0.001	47	36.6 (±13.7)	52	43.6 (±11.6)	0.006

Haemolysis score (0-6)																									
0	166	39.0%	0	0.0%	<0.001	112	41.2%	33	12.2%	<0.001	54	35.8%	17	32.7%	0.69	50	35.5%	0	0.0%	<0.001	17	32.7%	4	7.8%	0.001
1	144	34.0%	43	10.2%	<0.001	91	33.5%	135	49.9%	<0.001	53	35.1%	12	23.1%	0.11	51	36.2%	9	6.5%	<0.001	12	23.1%	21	41.2%	0.05
2	78	18.4%	213	50.6%	<0.001	49	18.00%	92	34.00%	<0.001	29	19.2%	16	30.7%	0.083	26	18.4%	75	54.0%	<0.001	16	30.8%	25	49.00%	0.06
3	35	8.3%	150	35.6%	<0.001	20	7.35%	11	4.1%	0.1	15	9.93%	7	13.5%	0.48	14	9.9%	51	36.7%	<0.001	7	13.5%	1	1.96%	0.03
4	0	0.0%	15	3.5%	<0.001	0	0.00%	0	0.00%	..	0	0.00%	0	0.00%	..	0	0.0%	4	2.9%	0.042	0	0.00%	0	0.00%	..

**Supplementary Table 5:** Further Iron, Infection and Haematological Parameters (Umbilical cord and venous sample) for FTB/NBW, PTB/LBW, PTB and LBW neonates. Data are presented as mean ( $\pm$  SD) and analysed by ANOVA. *P* values in bold font are considered significant based on  $P < 0.05$ .

**Supplementary Table 6 – Changes in iron status, inflammation and hepcidin between cord and post-natal blood (Groups A-D)**

	Whole Population					Group A (PTB+LBW)					Group B (PTB+NBW)					Group C (FTB+LBW)					Group D (FTB+NBW)				
Sample Type	n	Cord	n	Venous	P value	n	Cord	n	Venous	P value	n	Cord	n	Venous	P value	n	Cord	n	Venous	P value	n	Cord	n	Venous	P value
Serum iron (μmol/L)	425	23.3 (±7.1)	421	7.5 (±4.5)	<0.001	41	25.0 (±7.6)	40	8.2 (±3.3)	<0.001	99	24.2 (±7.2)	99	7.7 (±4.4)	<0.001	10	22.6 (±7.9)	11	9.4 (±3.9)	0.0039	275	22.7 (±7.0)	271	7.3 (±4.6)	<0.001
TSAT (%)	423	51.7 (±17.3)	420	15.0 (±6.9)	<0.001	41	55.6 (±17.7)	39	15.9 (±5.5)	<0.001	99	54.5 (±17.7)	99	16.2 (±9.3)	<0.001	10	50.4 (±23.2)	11	17.1 (±4.5)	0.0019	273	50.2 (±16.7)	271	14.4 (±6.1)	<0.001
Serum hepcidin (ng/ml)	425	19.9 (±14.2)	417	38.4 (±23.7)	<0.001	42	18.8 (±13.2)	40	33.6 (±21.1)	<0.001	96	22.7 (±14.1)	96	39.4 (±24.0)	<0.001	10	12.7 (±11.6)	11	34.0 (±27.0)	0.023	277	19.4 (±14.4)	270	38.9 (±23.9)	<0.001
UIBC (μmol/L)	423	22.9 (±10.5)	420	43.5 (±15.9)	<0.001	41	21.1 (±10.0)	39	43.4 (±10.1)	<0.001	99	21.2 (±9.8)	99	41.5 (±11.2)	<0.001	10	26.4 (±18.8)	11	45.2 (±10.9)	0.0006	273	23.7 (±10.4)	271	44.1 (±18.0)	<0.001
TIBC (μmol/L)	423	46.1 (±8.0)	420	51.0 (±17.9)	<0.001	41	46.0 (±8.4)	39	51.6 (±11.6)	0.0014	99	45.4 (±7.0)	99	49.1 (±10.9)	0.0013	10	49.0 (±12.6)	11	54.6 (±13.5)	0.022	273	46.3 (±8.1)	271	51.4 (±20.7)	<0.001
Serum ferritin (ug/L)	426	210.5 (±157.8)	415	381.2 (±297.6)	<0.001	42	189.3 (±173.1)	40	358.5 (±340.0)	0.0002	99	214.1 (±146.7)	98	354.6 (±220.7)	<0.001	10	203.8 (±214.4)	11	394.3 (±372.9)	0.05	275	212.6 (±157.7)	266	393.9 (±312.6)	<0.001
Haemoglobin (g/dl)	414	15.3 (±2.5)	423	19.4 (±3.1)	<0.001	40	16.4 (±2.9)	41	20.8 (±3.3)	<0.001	96	15.3 (±2.4)	99	19.4 (±3.1)	<0.001	8	15.8 (±4.5)	11	20.2 (±4.3)	0.021	270	15.1 (±2.3)	272	19.1 (±2.9)	<0.001
Hematocrit (%)	414	42.7 (±7.4)	423	54.7 (±9.3)	<0.001	40	46.1 (±8.3)	41	59.4 (±10.1)	<0.001	96	42.7 (±7.3)	99	54.5 (±9.4)	<0.001	8	44.7 (±13.3)	11	57.5 (±12.9)	0.023	270	42.2 (±7.0)	272	53.9 (±8.9)	<0.001
Transferrin (g/L)	426	1.96 (±0.3)	423	1.99 (±0.3)	0.003	42	1.90 (±0.3)	40	2.00 (±0.4)	0.0034	99	1.92 (±0.3)	99	1.99 (±0.3)	0.0001	10	2.08 (±0.6)	11	2.11 (±0.5)	0.43	275	1.98 (±0.3)	273	1.98 (±0.3)	0.84
Soluble transferrin receptor (mg/L)	424	6.0 (±2.0)	421	7.0 (±2.2)	<0.001	42	6.14 (±2.0)	40	7.47 (±2.3)	<0.001	99	5.74 (±1.6)	99	6.67 (±1.8)	<0.001	10	7.00 (±3.7)	11	8.6 (±2.9)	0.0011	273	5.99 (±2.0)	271	6.94 (±2.2)	<0.001
Haptoglobin (g/L)	426	0.02 (±0.07)	423	0.02 (±0.06)	0.52	42	0.01 (±0.01)	40	0.01 (±0.01)	0.18	99	0.03 (±0.1)	99	0.02 (±0.05)	0.15	10	0.02 (±0.03)	11	0.02 (±0.02)	0.87	275	0.02 (±0.06)	273	0.03 (±0.07)	0.06
Undetectable haptoglobin (%)	182	42.7%	144	34.0%	0.009	20	47.6%	10	25.00%	0.033	42	42.4%	28	28.3%	0.037	4	40%	5	45.5%	0.8	116	42.2%	101	37.00%	0.21
Serum AGP (g/L)	426	0.18 (±0.1)	423	0.24 (±0.2)	<0.001	42	0.15 (±0.2)	40	0.19 (±0.2)	<0.001	99	0.17 (±0.2)	99	0.24 (±0.2)	<0.001	10	0.22 (±0.1)	11	0.27 (±0.15)	0.063	275	0.19 (±0.1)	273	0.25 (±0.17)	<0.001
Serum CRP (mg/L)	425	0.17 (±0.6)	422	2.16 (±4.0)	<0.001	42	0.09 (±0.05)	40	1.01 (±1.2)	<0.001	99	0.16 (±0.3)	98	2.39 (±4.5)	<0.001	10	0.14 (±0.1)	11	1.63 (±3.3)	0.19	275	0.19 (±0.7)	273	2.27 (±4.1)	<0.001
>5mg/L	2	0.5%	54	12.8%	<0.001	0	0.0%	3	7.5%	N/A	0	0.0%	13	13.3%	N/A	0	0.0%	1	9.1%	N/A	2	0.7%	37	13.6%	<0.001
>3mg/L	3	0.7%	83	19.7%	<0.001	0	0.0%	5	12.5%	N/A	0	0.0%	18	18.4%	N/A	0	0.0%	1	9.1%	N/A	3	1.0%	59	21.6%	<0.001
Mean corpuscular volume (fl)	414	98.0 (±6.2)	423	98.0 (±6.1)	0.63	40	100.0 (±6.1)	41	100.1 (±6.5)	0.17	96	98.2 (±5.4)	99	98.3 (±5.4)	0.76	8	95.3 (±11.4)	11	94.2 (±9.8)	0.28	270	97.8 (±6.3)	272	97.7 (±6.0)	0.29
Mean corpuscular haemoglobin (pg)	414	35.1 (±2.4)	423	34.9 (±2.4)	<0.001	40	35.6 (±2.6)	41	35.2 (±2.5)	0.0001	96	35.3 (±2.2)	99	35.1 (±2.1)	0.0001	8	33.9 (±4.3)	11	33.2 (±3.8)	0.46	270	35.1 (±2.4)	272	34.8 (±2.4)	<0.001
Mean corpuscular haemoglobin concentration (g/dl)	414	35.8 (±1.0)	423	35.6 (±1.0)	<0.001	40	35.5 (±0.9)	41	35.2 (±0.9)	0.0034	96	35.9 (±1.0)	99	35.7 (±0.9)	0.0031	8	35.5 (±1.1)	11	35.3 (±0.9)	0.62	270	35.8 (±0.9)	272	35.6 (±1.1)	<0.001
White blood cell count (unit/L)	412	13.0 (±5.3)	422	15.9 (±5.5)	<0.001	40	10.9 (±4.2)	41	13.1 (±4.2)	0.0027	95	12.7 (±4.4)	99	15.8 (±4.8)	<0.001	8	14.7 (±6.6)	11	14.0 (±3.7)	0.8	269	13.3 (±5.7)	271	16.4 (±5.8)	<0.001
Lymphocyte count (unit/L)	412	4.73 (±3.1)	422	4.1 (±2.4)	<0.001	40	4.5 (±2.0)	41	3.7 (±1.2)	0.021	95	4.6 (±2.0)	99	4.0 (±1.5)	0.01	8	6.6 (±3.6)	11	3.6 (±1.1)	0.052	269	4.77 (±3.5)	271	4.21 (±2.9)	0.0001
Lymphocyte percentage (%)	412	36.2 (±8.3)	422	26.9 (±8.8)	<0.001	40	41.8 (±9.3)	41	30.5 (±9.6)	<0.001	95	36.2 (±7.9)	99	26.8 (±8.5)	<0.001	8	43.4 (±6.4)	11	27.0 (±7.2)	0.0048	269	35.2 (±7.9)	271	26.5 (±8.8)	<0.001
MID cell count (unit/L)	412	1.02 (±0.6)	422	1.48 (±1.0)	<0.001	40	1.01 (±0.6)	41	1.25 (±0.9)	0.13	95	0.97 (±0.4)	99	1.42 (±1.0)	<0.001	8	1.23 (±0.5)	11	1.32 (±1.0)	0.5	269	1.05 (±0.6)	271	1.55 (±1.0)	<0.001
MID cell percentage (%)	412	8.27 (±2.6)	422	9.67 (±5.4)	<0.001	40	9.56 (±4.4)	41	9.79 (±5.2)	0.75	95	8.04 (±2.0)	99	9.27 (±4.9)	0.017	8	8.99 (±3.5)	11	9.62 (±5.3)	0.61	269	8.15 (±2.4)	271	9.81 (±5.6)	<0.001

Granulocyte count (unit/L)	412	7.21 (±2.7)	422	10.3 (±4.1)	<0.001	40	5.43 (±2.6)	41	8.12 (±3.4)	<0.001	95	7.18 (±2.7)	99	10.32 (±3.9)	<0.001	8	6.9 (±3.0)	11	9.02 (±2.8)	0.046	269	7.49 (±2.7)	271	10.67 (±4.2)	<0.001	
Granulocyte percentage (%)	412	55.5 (±9.1)	422	63.4 (±8.6)	<0.001	40	48.7 (±10.2)	41	59.8 (±8.5)	<0.001	95	55.8 (±8.2)	99	64.0 (±7.8)	<0.001	8	47.6 (±7.8)	11	63.3 (±6.6)	0.0013	269	56.6 (±8.7)	271	63.7 (±8.8)	<0.001	
Red blood cell count (unit/L)	414	4.36 (±0.7)	423	5.58 (±1.0)	<0.001	40	4.6 (±0.8)	41	5.9 (±1.0)	<0.001	96	4.4 (±0.7)	99	5.5 (±0.9)	<0.001	8	4.73 (±1.3)	11	6.11 (±1.2)	0.016	270	4.32 (±0.7)	272	5.53 (±0.9)	<0.001	
Red blood cell distribution width (%)	414	15.2 (±1.1)	423	15.4 (±1.3)	<0.001	40	15.1 (±0.8)	41	15.2 (±0.8)	0.0069	96	15.2 (±1.0)	99	15.4 (±1.0)	0.0001	8	15.8 (±1.3)	11	15.6 (±1.1)	0.096	270	15.2 (±1.2)	272	15.4 (±1.4)	<0.001	
Red blood cell distribution width - absolute (fl)	414	81.0 (±8.6)	423	82.2 (±8.5)	<0.001	40	82.4 (±7.8)	41	83.8 (±8.4)	0.0038	96	81.2 (±7.7)	99	82.5 (±7.9)	0.0008	8	79.8 (±12.7)	11	79.2 (±11.5)	0.38	270	80.7 (±8.9)	272	82.0 (±8.6)	0.0001	
Platelet count (unit/L)	414	241.4 (±86.9)	423	260.4 (±94.6)	0.0003	40	226.3 (±76.4)	41	256.9 (±86.1)	0.092	96	242.2 (±70.1)	99	269.4 (±101.0)	0.015	8	223.8 (±117.5)	11	234.3 (±84.3)	0.76	270	243.8 (±92.8)	272	258.7 (±94.0)	0.02	
Mean platelet volume (fl)	409	8.34 (±0.8)	419	8.47 (±0.8)	0.0003	39	8.16 (±0.8)	41	8.35 (±0.8)	0.083	95	8.36 (±0.8)	96	8.38 (±0.7)	0.99	8	8.22 (±0.8)	11	8.46 (±0.8)	0.5	267	8.37 (±0.8)	271	8.52 (±0.8)	0.0001	
Platelet distribution width (%)	409	43.3 (±2.8)	419	43.7 (±3.2)	0.007	39	43.0 (±2.9)	41	43.2 (±3.1)	0.94	95	42.9 (±2.4)	96	43.4 (±2.8)	0.17	8	42.6 (±2.4)	11	44.2 (±2.7)	0.0064	267	43.5 (±3.0)	271	43.9 (±3.3)	0.021	
Platelet distribution width - absolute (fl)	409	11.8 (±1.2)	419	12.0 (±1.3)	0.0006	39	11.5 (±1.1)	41	11.7 (±1.3)	0.33	95	11.7 (±1.2)	96	11.8 (±1.1)	0.56	8	11.6 (±1.2)	11	12.1 (±1.3)	0.026	267	11.8 (±1.3)	271	12.1 (±1.4)	0.0009	
Platelet crit (%)	409	0.2 (±0.06)	419	0.22 (±0.07)	<0.001	39	0.183 (±0.05)	41	0.212 (±0.06)	0.031	95	0.199 (±0.06)	96	0.226 (±0.07)	0.0008	8	0.179 (±0.09)	11	0.190 (±0.07)	0.87	267	0.201 (±0.07)	271	0.215 (±0.07)	0.01	
Plate large cell ratio (%)	409	17.8 (±5.3)	419	18.8 (±5.5)	0.0001	39	16.6 (±5.0)	41	17.9 (±5.3)	0.1	95	17.8 (±5.2)	96	18.1 (±4.7)	0.86	8	17.6 (±4.9)	11	19.3 (±5.4)	0.062	267	18.0 (±5.4)	271	19.2 (±5.8)	0.0001	
Plate large cell count (unit/L)	409	41.4 (±14.2)	419	46.5 (±15.8)	<0.001	39	36.5 (±12.2)	41	44.1 (±11.3)	0.005	95	41.7 (±12.6)	96	47.2 (±14.1)	0.0025	8	37.3 (±20.7)	11	41.8 (±13.0)	0.71	267	42.1 (±14.6)	271	46.9 (±17.0)	<0.001	
Haemolysis score (0-6)	0	166	39.0%	0	0.0%	<0.001	13	31.00%	3	7.5%	0.007	37	37.4%	6	6.1%	<0.001	4	40.00%	0	0.00%	0.019	112	41.2%	33	12.2%	<0.001
	1	144	34.0%	43	10.2%	<0.001	10	23.8%	18	45.00%	0.043	41	41.4%	57	57.6%	0.022	2	20.00%	1	9.1%	0.48	91	33.5%	135	49.8%	<0.001
	2	78	18.4%	213	50.6%	<0.001	13	31.00%	18	45.00%	0.18	13	13.1%	33	33.3%	<0.001	3	30.00%	3	27.3%	0.89	49	18.00%	92	34.00%	<0.001
	3	35	8.3%	150	35.6%	<0.001	6	14.3%	1	4.5%	0.056	8	8.1%	3	3.00%	0.12	1	10.00%	7	63.6%	0.011	20	7.4%	11	4.1%	0.098
	4	0	0.0%	15	3.5%	<0.001	0	0.00%	0	0.00%	..	0	0.00%	0	0.00%	..	0	0.00%	0	0.00%	..	0	0.00%	0	0.00%	..

**Supplementary Table 6:** Iron, Infection and Haematological Parameters (umbilical cord and venous samples) for groups A, B, C and D

neonates. Data are presented as mean (± SD) and analysed by ANOVA. *P* values in bold font are considered significant based on *P*<0.05.

**Supplementary Table 7 – Comparison of iron status, inflammation and haematological parameters in cord samples (Group A-D)**

	Whole Population		Group A (PTB+LBW)		Group B (PTB+NBW)		Group C (FTB+LBW)		Group D (FTB+NBW)		
Sample Type	n	Cord	n	Cord	n	Cord	n	Cord	n	Cord	P value
Serum iron (μmol/L)	425	23·3 (±7·1)	41	25·0 (±7·6)	99	24·2 (±7·2)	10	22·6 (±7·9)	275	22·7 (±7·0)	0·11
TSAT (%)	423	51·7 (±17·3)	41	55·6 (±17·7)	99	54·5 (±17·7)	10	50·4 (±23·2)	273	50·2 (±16·7)	0·079
Serum hepcidin (ng/ml)	425	19·9 (±14·2)	42	18·8 (±13·2)	96	22·7 (±14·1)	10	12·7 (±11·6)	277	19·4 (±14·4)	0·075
UIBC (μmol/L)	423	22·9 (±10·5)	41	21·1 (±10·0)	99	21·2 (±9·8)	10	26·4 (±18·8)	273	23·7 (±10·4)	0·094
TIBC (μmol/L)	423	46·1 (±8·0)	41	46·0 (±8·4)	99	45·4 (±7·0)	10	49·0 (±12·6)	273	46·3 (±8·1)	0·52
Serum ferritin (ug/L)	426	210·5 (±157·8)	42	189·3 (±173·1)	99	214·1 (±146·7)	10	203·8 (±214·4)	275	212·6 (±157·7)	0·83
Haemoglobin (g/dl)	414	15·3 (±2·5)	40	16·4 (±2·9)	96	15·3 (±2·4)	8	15·8 (±4·5)	270	15·1 (±2·3)	<b>0·02</b>
Haematocrit (%)	414	42·7 (±7·4)	40	46·1 (±8·3)	96	42·7 (±7·3)	8	44·7 (±13·3)	270	42·2 (±7·0)	<b>0·015</b>
Transferrin (g/L)	426	1·96 (±0·3)	42	1·90 (±0·3)	99	1·92 (±0·3)	10	2·08 (±0·6)	275	1·98 (±0·3)	0·18
Soluble transferrin receptor (mg/L)	424	6·0 (±2·0)	42	6·14 (±2·0)	99	5·74 (±1·6)	10	7·00 (±3·7)	273	5·99 (±2·0)	0·22
Haptoglobin (g/L)	426	0·02 (±0·07)	42	0·01 (±0·01)	99	0·03 (±0·1)	10	0·02 (±0·03)	275	0·02 (±0·06)	0·21
Undetectable haptoglobin (%)	182	42·7%	20	47·6%	42	42·4%	4	40%	116	42·2%	0·92
Serum AGP (g/L)	426	0·18 (±0·1)	42	0·15 (±0·2)	99	0·17 (±0·2)	10	0·22 (±0·1)	275	0·19 (±0·1)	0·24
Serum CRP (mg/L)	425	0·17 (±0·6)	42	0·09 (±0·05)	99	0·16 (±0·3)	10	0·14 (±0·1)	275	0·19 (±0·7)	0·75
>5mg/L	2	0·5%	0	0·0%	0	0·0%	0	0·0%	2	0·7%	0·78
>3mg/L	3	0·7%	0	0·0%	0	0·0%	0	0·0%	3	1·0%	0·65
Mean corpuscular volume (fl)	414	98·0 (±6·2)	40	100·0 (±6·1)	96	98·2 (±5·4)	8	95·3 (±11·4)	270	97·8 (±6·3)	0·1
Mean corpuscular haemoglobin (pg)	414	35·1 (±2·4)	40	35·6 (±2·6)	96	35·3 (±2·2)	8	33·9 (±4·3)	270	35·1 (±2·4)	0·25
Mean corpuscular haemoglobin concentration (g/dl)	414	35·8 (±1·0)	40	35·5 (±0·9)	96	35·9 (±1·0)	8	35·5 (±1·1)	270	35·8 (±0·9)	0·17
White blood cell count (unit/L)	412	13·0 (±5·3)	40	10·9 (±4·2)	95	12·7 (±4·4)	8	14·7 (±6·6)	269	13·3 (±5·7)	<b>0·04</b>
Lymphocyte count (unit/L)	412	4·73 (±3·1)	40	4·5 (±2·0)	95	4·6 (±2·0)	8	6·6 (±3·6)	269	4·77 (±3·5)	0·32
Lymphocyte percentage (%)	412	36·2 (±8·3)	40	41·8 (±9·3)	95	36·2 (±7·9)	8	43·4 (±6·4)	269	35·2 (±7·9)	<b>&lt;0·001</b>
MID cell count (unit/L)	412	1·02 (±0·6)	40	1·01 (±0·6)	95	0·97 (±0·4)	8	1·23 (±0·5)	269	1·05 (±0·6)	0·49
MID cell percentage (%)	412	8·27 (±2·6)	40	9·56 (±4·4)	95	8·04 (±2·0)	8	8·99 (±3·5)	269	8·15 (±2·4)	<b>0·009</b>



Granulocyte count (unit/L)	412	7.21 (±2.7)	40	5.43 (±2.6)	95	7.18 (±2.7)	8	6.9 (±3.0)	269	7.49 (±2.7)	0.0002	
Granulocyte percentage (%)	412	55.5 (±9.1)	40	48.7 (±10.2)	95	55.8 (±8.2)	8	47.6 (±7.8)	269	56.6 (±8.7)	<0.001	
Red blood cell count (unit/L)	414	4.36 (±0.7)	40	4.6 (±0.8)	96	4.4 (±0.7)	8	4.73 (±1.3)	270	4.32 (±0.7)	0.054	
Red blood cell distribution width (%)	414	15.2 (±1.1)	40	15.1 (±0.8)	96	15.2 (±1.0)	8	15.8 (±1.3)	270	15.2 (±1.2)	0.45	
Red blood cell distribution width - absolute (fl)	414	81.0 (±8.6)	40	82.4 (±7.8)	96	81.2 (±7.7)	8	79.8 (±12.7)	270	80.7 (±8.9)	0.67	
Platelet count (unit/L)	414	241.4 (±86.9)	40	226.3 (±76.4)	96	242.2 (±70.1)	8	223.8 (±117.5)	270	243.8 (±92.8)	0.63	
Mean platelet volume (fl)	409	8.34 (±0.8)	39	8.16 (±0.8)	95	8.36 (±0.8)	8	8.22 (±0.8)	267	8.37 (±0.8)	0.41	
Platelet distribution width (%)	409	43.3 (±2.8)	39	43.0 (±2.9)	95	42.9 (±2.4)	8	42.6 (±2.4)	267	43.5 (±3.0)	0.31	
Platelet distribution width - absolute (fl)	409	11.8 (±1.2)	39	11.5 (±1.1)	95	11.7 (±1.2)	8	11.6 (±1.2)	267	11.8 (±1.3)	0.39	
Platelet crit (%)	409	0.2 (±0.06)	39	0.183 (±0.05)	95	0.199 (±0.06)	8	0.179 (±0.09)	267	0.201 (±0.07)	0.29	
Plate large cell ratio (%)	409	17.8 (±5.3)	39	16.6 (±5.0)	95	17.8 (±5.2)	8	17.6 (±4.9)	267	18.0 (±5.4)	0.47	
Plate large cell count (unit/L)	409	41.4 (±14.2)	39	36.5 (±12.2)	95	41.7 (±12.6)	8	37.3 (±20.7)	267	42.1 (±14.6)	0.11	
Haemolysis score (0-6)												
	0	166	39.0%	13	31.00%	37	37.4%	4	40.00%	112	41.2%	0.62
	1	144	34.0%	10	23.8%	41	41.4%	2	20.00%	91	33.5%	0.2
	2	78	18.4%	13	31.00%	13	13.1%	3	30.00%	49	18.00%	0.067
	3	35	8.3%	6	14.3%	8	8.1%	1	10.00%	20	7.4%	0.5
	4	0	0.0%	0	0.00%	0	0.00%	0	0.00%	0	0.00%	..

**Supplementary Table 7:** Iron, infection and haematological parameters (umbilical cord samples) for groups A, B, C, and D neonates. Data are presented as mean (± SD) and analysed by ANOVA. *P* values in bold font are considered significant based on *P*<0.05.

**Supplementary Table 8 – Comparison of iron status, inflammation and haematological parameters in venous samples (Groups A-D)**

	Whole Population		Group A (PTB+LBW)		Group B (PTB+NBW)		Group C (FTB+LBW)		Group D (FTB+NBW)		
Sample Type	n	Venous	n	Venous	n	Venous	n	Venous	n	Venous	P value
Serum iron (μmol/L)	421	7.5 (±4.5)	40	8.2 (±3.3)	99	7.7 (±4.4)	11	9.4 (±3.9)	271	7.3 (±4.6)	0.29
TSAT (%)	420	15.0 (±6.9)	39	15.9 (±5.5)	99	16.2 (±9.3)	11	17.1 (±4.5)	271	14.4 (±6.1)	0.68
Serum hepcidin (ng/ml)	417	38.4 (±23.7)	40	33.6 (±21.1)	96	39.4 (±24.0)	11	34.0 (±27.0)	270	38.9 (±23.9)	0.076
UIBC (μmol/L)	420	43.5 (±15.9)	39	43.4 (±10.1)	99	41.5 (±11.2)	11	45.2 (±10.9)	271	44.1 (±18.0)	0.53
TIBC (μmol/L)	420	51.0 (±17.9)	39	51.6 (±11.6)	99	49.1 (±10.9)	11	54.6 (±13.5)	271	51.4 (±20.7)	0.64
Serum ferritin (ug/L)	415	381.2 (±297.6)	40	358.5 (±340.0)	98	354.6 (±220.7)	11	394.3 (±372.9)	266	393.9 (±312.6)	0.68
Haemoglobin (g/dl)	423	19.4 (±3.1)	41	20.8 (±3.3)	99	19.4 (±3.1)	11	20.2 (±4.3)	272	19.1 (±2.9)	<b>0.009</b>
Haematocrit (%)	423	54.7 (±9.3)	41	59.4 (±10.1)	99	54.5 (±9.4)	11	57.5 (±12.9)	272	53.9 (±8.9)	<b>0.0042</b>
Transferrin (g/L)	423	1.99 (±0.3)	40	2.00 (±0.4)	99	1.99 (±0.3)	11	2.11 (±0.5)	273	1.98 (±0.3)	0.65
Soluble transferrin receptor (mg/L)	421	7.0 (±2.2)	40	7.47 (±2.3)	99	6.67 (±1.8)	11	8.6 (±2.9)	271	6.94 (±2.2)	<b>0.017</b>
Haptoglobin (g/L)	423	0.02 (±0.06)	40	0.01 (±0.01)	99	0.02 (±0.05)	11	0.02 (±0.02)	273	0.03 (±0.07)	0.67
Undetectable haptoglobin (%)	144	34.0%	10	25.00%	28	28.3%	5	45.5%	101	37.00%	0.2
Serum AGP (g/L)	423	0.24 (±0.2)	40	0.19 (±0.2)	99	0.24 (±0.2)	11	0.27 (±0.15)	273	0.25 (±0.17)	0.19
Serum CRP (mg/L)	422	2.16 (±4.0)	40	1.01 (±1.2)	98	2.39 (±4.5)	11	1.63 (±3.3)	273	2.27 (±4.1)	0.26
>5mg/L %	54	12.8%	3	7.5%	13	13.3%	1	9.1%	37	13.6%	0.73
>3mg/L %	83	19.7%	5	12.5%	18	18.4%	1	9.1%	59	21.6%	0.42
Mean corpuscular volume (fl)	423	98.0 (±6.1)	41	100.1 (±6.5)	99	98.3 (±5.4)	11	94.2 (±9.8)	272	97.7 (±6.0)	<b>0.017</b>
Mean corpuscular haemoglobin (pg)	423	34.9 (±2.4)	41	35.2 (±2.5)	99	35.1 (±2.1)	11	33.2 (±3.8)	272	34.8 (±2.4)	0.08
Mean corpuscular haemoglobin concentration (g/dl)	423	35.6 (±1.0)	41	35.2 (±0.9)	99	35.7 (±0.9)	11	35.3 (±0.9)	272	35.6 (±1.1)	<b>0.029</b>
White blood cell count (unit/L)	422	15.9 (±5.5)	41	13.1 (±4.2)	99	15.8 (±4.8)	11	14.0 (±3.7)	271	16.4 (±5.8)	<b>0.0021</b>
Lymphocyte count (unit/L)	422	4.1 (±2.4)	41	3.7 (±1.2)	99	4.0 (±1.5)	11	3.6 (±1.1)	271	4.21 (±2.9)	0.6
Lymphocyte percentage (%)	422	26.9 (±8.8)	41	30.5 (±9.6)	99	26.8 (±8.5)	11	27.0 (±7.2)	271	26.5 (±8.8)	0.06
MID cell count (unit/L)	422	1.48 (±1.0)	41	1.25 (±0.9)	99	1.42 (±1.0)	11	1.32 (±1.0)	271	1.55 (±1.0)	0.27
MID cell percentage (%)	422	9.67 (±5.4)	41	9.79 (±5.2)	99	9.27 (±4.9)	11	9.62 (±5.3)	271	9.81 (±5.6)	0.86

Granulocyte count (unit/L)	422	10.3 (±4.1)	41	8.12 (±3.4)	99	10.32 (±3.9)	11	9.02 (±2.8)	271	10.67 (±4.2)	0.0017	
Granulocyte percentage (%)	422	63.4 (±8.6)	41	59.8 (±8.5)	99	64.0 (±7.8)	11	63.3 (±6.6)	271	63.7 (±8.8)	0.04	
Red blood cell count (unit/L)	423	5.58 (±1.0)	41	5.9 (±1.0)	99	5.5 (±0.9)	11	6.11 (±1.2)	272	5.53 (±0.9)	0.018	
Red blood cell distribution width (%)	423	15.4 (±1.3)	41	15.2 (±0.8)	99	15.4 (±1.0)	11	15.6 (±1.1)	272	15.4 (±1.4)	0.69	
Red blood cell distribution width - absolute (fl)	423	82.2 (±8.5)	41	83.8 (±8.4)	99	82.5 (±7.9)	11	79.2 (±11.5)	272	82.0 (±8.6)	0.39	
Platelet count (unit/L)	423	260.4 (±94.6)	41	256.9 (±86.1)	99	269.4 (±101.0)	11	234.3 (±84.3)	272	258.7 (±94.0)	0.35	
Mean platelet volume (fl)	419	8.47 (±0.8)	41	8.35 (±0.8)	96	8.38 (±0.7)	11	8.46 (±0.8)	271	8.52 (±0.8)	0.3	
Platelet distribution width (%)	419	43.7 (±3.2)	41	43.2 (±3.1)	96	43.4 (±2.8)	11	44.2 (±2.7)	271	43.9 (±3.3)	0.26	
Platelet distribution width - absolute (fl)	419	12.0 (±1.3)	41	11.7 (±1.3)	96	11.8 (±1.1)	11	12.1 (±1.3)	271	12.1 (±1.4)	0.19	
Platelet crit (%)	419	0.22 (±0.07)	41	0.212 (±0.06)	96	0.226 (±0.07)	11	0.190 (±0.07)	271	0.215 (±0.07)	0.35	
Plate large cell ratio (%)	419	18.8 (±5.5)	41	17.9 (±5.3)	96	18.1 (±4.7)	11	19.3 (±5.4)	271	19.2 (±5.8)	0.21	
Plate large cell count (unit/L)	419	46.5 (±15.8)	41	44.1 (±11.3)	96	47.2 (±14.1)	11	41.8 (±13.0)	271	46.9 (±17.0)	0.52	
Haemolysis score (0-6)												
	0	0	0.0%	3	7.5%	6	6.1%	0	0.00%	33	12.2%	0.2
	1	43	10.2%	18	45.00%	57	57.6%	1	9.1%	135	49.8%	0.018
	2	213	50.6%	18	45.00%	33	33.3%	3	27.3%	92	34.00%	0.51
	3	150	35.6%	1	4.5%	3	3.00%	7	63.6%	11	4.1%	<0.001
	4	15	3.5%	0	0.00%	0	0.00%	0	0.00%	0	0.00%	..

**Supplementary Table 8:** Iron, infection and haematological parameters (venous samples) for groups A, B, C, and D neonates. Data are presented as mean (± SD) and analysed by ANOVA. *P* values in bold font are considered significant based on *P* < 0.05.

## SUPPLEMENTARY REFERENCES

- 1 Cross JH, Jarjou O, Mohammed NI, Prentice AM, Cerami C. Neonatal iron distribution and infection susceptibility in full term, preterm and low birthweight babies in urban Gambia: study protocol for an observational study. *Gates Open Res* 2019; **3**: 1469.

## Chapter 6 – Iron homeostasis in full term, normal birthweight Gambian neonates over the first week of life (FTB/NBW Paper)

---

### Summary of Chapter

**BACKGROUND:** Human neonates elicit a profound hypoferremia to protect against bacterial and fungal sepsis on their first day of life.

**OBJECTIVE:** We examined the transience of this effect by measuring iron and its chaperone proteins, inflammatory and hematological parameters over the first post-partum week.

**DESIGN:** We prospectively studied term (>37 completed gestational weeks), normal weight (>2500g) newborns at Kanifing General Hospital, The Gambia. Blood was sampled from the umbilical cord vein (CDV) and artery (CDA). Neonatal venous blood was sampled at 6-24h (V1) in all babies, who were then randomized to a second blood draw at 25-80h (V2), 81-136h (V3) or 137-192h (V4). Hepcidin, serum iron, transferrin saturation, transferrin, haptoglobin, CRP, AGP, sTfR, ferritin, TIBC, UIBC and full blood count were assayed.

**RESULTS:** Two hundred and seventy-eight neonates (278, 54.3% males, gestational age  $39.4 \pm 1.3$  wk, birth weight  $3299 \pm 368$  g) were enrolled. We confirmed the profound early postnatal decrease in serum iron (CDV =  $22.7 \pm 7.0$   $\mu$ mol/L to V1 =  $7.3 \pm 4.3$   $\mu$ mol/L,  $P < 0.0001$ ) and TSAT ( $50.2 \pm 16.7\%$  to  $14.4 \pm 6.1\%$ ,  $P < 0.0001$ ). Both variables increased steadily to reach  $16.5 \pm 3.9$   $\mu$ mol/L and  $36.7 \pm 9.2\%$  at V4 ( $P$  for trend  $< 0.0001$  for each). Hepcidin increased rapidly after birth (CDV =  $19.4 \pm 14.4$  ng/ml to V1 =  $38.9 \pm 23.9$  ng/ml,  $P < 0.0001$ ) then dipped before rising again by V4. Inflammatory markers increased from V1 onwards. Network

analysis revealed a disconnect between the correlations observed in CDV and V2-V4 and those in V1. Surprisingly, serum iron and TSAT were only weakly influenced by hepcidin.

**CONCLUSIONS:** Rapid anti-infective post-natal hypoferremia in human neonates is transient. The later rise in serum iron despite very high hepcidin indicates hepcidin resistance possibly caused by macrophage saturation with iron released from degradation of fetal erythrocytes. Pharmacological prolongation of hypoferremia might offer an ancillary tool in the armoury against antimicrobial resistance, however, it would need to overcome or circumvent the hepcidin resistance we now report.

## RESEARCH PAPER COVER SHEET - FTB/NBW PAPER

---

Please note that a cover sheet must be completed for each research paper included within a thesis.

### **SECTION A – Student Details**

<b>Student ID Number</b>	LSH158152	<b>Title</b>	Mr.
<b>First Name(s)</b>	James Henry		
<b>Surname/Family Name</b>	Cross		
<b>Thesis Title</b>	Iron and Infection: Neonatal Iron Transition		
<b>Primary Supervisor</b>	Dr Carla Cerami		

If the Research Paper has previously been published please complete Section B, if not please move to Section C.

### **SECTION B – Paper already published**

Where was the work published?

When was the work published?

If the work was published prior to registration for your research degree, give a brief rationale for its inclusion

Have you retained the copyright for the work?*	Choose an item.	Was the work subject to academic peer review?	Choose an item.
--	-----------------	---	-----------------

\*If yes, please attach evidence of retention. If no, or if the work is being included in its published format, please attach evidence of permission from the copyright holder (publisher or other author) to include this work.

### **SECTION C – Prepared for publication, but not yet published**

Where is the work intended to be published?	American Journal of Clinical Nutrition
Please list the paper's authors in the intended authorship order:	James H. Cross, Ousman Jarjou, Nuredin Ibrahim Mohammed, Santiago Rayment Gomez, Bubacarr J.B Touray, Andrew M. Prentice, Carla Cerami.
Stage of publication	<b>Undergoing revision</b>

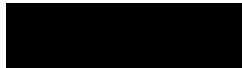
## **SECTION D – Multi-authored work**

For multi-authored work, give full details of your role in the research included in the paper and in the preparation of the paper. (Attach a further sheet if necessary)

James H. Cross is PhD student with MRCG at LSHTM. He contributed to the development of the protocols, study management, field data collection, development of the data analysis plan, conducting the data analysis and drafting of the manuscript.

## **SECTION E**

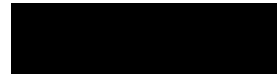
**Student Signature**



**Date**

08/11/2019

**Supervisor Signature**



**Date**

08/11/2019



**Title**

**Iron homeostasis in full term, normal birthweight Gambian neonates over the first week of life.**

**Authors**

James H. Cross<sup>1</sup>, Ousman Jarjou<sup>1</sup>, Nuredin Ibrahim Mohammed<sup>1</sup>, Santiago Rayment Gomez<sup>2</sup>, Bubacarr J.B Touray<sup>1</sup>, Andrew M. Prentice<sup>1</sup>, Carla Cerami<sup>1\*</sup>.

**Affiliations**

<sup>1</sup>Medical Research Council Unit The Gambia at the London School of Hygiene & Tropical Medicine, Atlantic Boulevard, Fajara, P.O. Box 273, Banjul, The Gambia

<sup>2</sup>Cardiff University, Cardiff, Wales, CF10 3AT, United Kingdom

**Corresponding Author**

Dr Carla Cerami, email: [ccerami@mrc.gm](mailto:ccerami@mrc.gm)

Medical Research Council Unit The Gambia at the London School of Hygiene & Tropical Medicine, Atlantic Boulevard, Fajara, P.O. Box 273, Banjul, The Gambia

+220 787 5756

**Running Title:**

Neonatal hypoferrremia in full term, normal birthweight neonates

## 6.1 ABSTRACT

**BACKGROUND:** Human neonates elicit a profound hypoferremia to protect against bacterial and fungal sepsis on their first day of life.

**OBJECTIVE:** We examined the transience of this effect by measuring iron and its chaperone proteins, inflammatory and hematological parameters over the first postpartum week.

**DESIGN:** We prospectively studied term (>37 completed gestational weeks), normal weight (>2500g) newborns at Kanifing General Hospital, The Gambia. Blood was sampled from the umbilical cord vein (CDV) and artery (CDA). Neonatal venous blood was sampled at 6-24h (V1) in all babies, who were then randomized to a second blood draw at 25-80h (V2), 81-136h (V3) or 137-192h (V4). Hepcidin, serum iron, transferrin saturation, transferrin, haptoglobin, CRP, AGP, sTfR, ferritin, TIBC, UIBC and full blood count were assayed.

**RESULTS:** Two hundred and seventy-eight neonates (278, 54.3% males, gestational age  $39.4 \pm 1.3$  wk, birth weight  $3299 \pm 368$  g) were enrolled. We confirmed the profound early postnatal decrease in serum iron (CDV =  $22.7 \pm 7.0$   $\mu$ mol/L to V1 =  $7.3 \pm 4.3$   $\mu$ mol/L,  $P < 0.0001$ ) and TSAT ( $50.2 \pm 16.7\%$  to  $14.4 \pm 6.1\%$ ,  $P < 0.0001$ ). Both variables increased steadily to reach  $16.5 \pm 3.9$   $\mu$ mol/L and  $36.7 \pm 9.2\%$  at V4 ( $P$  for trend  $< 0.0001$  for each). Hepcidin increased rapidly after birth (CDV =  $19.4 \pm 14.4$  ng/ml to V1 =  $38.9 \pm 23.9$  ng/ml,  $P < 0.0001$ ) then dipped before rising again by V4. Inflammatory markers increased from V1 onwards. Network analysis revealed a disconnect between the correlations observed in CDV and V2-V4 and those in V1. Surprisingly, serum iron and TSAT were only weakly influenced by hepcidin.

**CONCLUSIONS:** Rapid anti-infective postnatal hypoferremia in human neonates is transient. The later rise in serum iron despite very high hepcidin indicates hepcidin resistance possibly caused by macrophage saturation with iron released from degradation of fetal erythrocytes. Pharmacological prolongation of hypoferremia might offer an ancillary tool

in the armoury against antimicrobial resistance, however, it would need to overcome or circumvent the hepcidin resistance we now report.

**Trial Registration:** [clinicaltrials.gov](https://clinicaltrials.gov) (NCT03353051). Registration date: November 27, 2017.

**FUNDING:** Bill & Melinda Gates Foundation (OPP1152353).

**WORD COUNT:** 296/300

**Keywords:** Nutritional immunity, host-pathogen interaction, hepcidin, neonates, hypoferremia, transferrin, The Gambia, sub-Saharan Africa

## 6.2 INTRODUCTION

During pregnancy, the mother increases iron absorption and turnover of erythrocytes to provide for the growing fetus.<sup>1</sup> As maternal hepcidin decreases during the third trimester, placental iron transfer rises.<sup>2</sup> This leads to higher cord blood TSAT and serum iron levels compared to those of the mother at delivery,<sup>3–5</sup> even in anemic mothers.<sup>6</sup> To protect the fetus against possible iron overload during the last trimester, fetal-derived hepcidin regulates iron transfer via degradation of ferroportin on placental syncytiotrophoblasts.<sup>7</sup> As a result, umbilical cord hepcidin concentrations of term neonates are higher than those of the mother, before and during delivery.<sup>8–10</sup>

Using murine and *in vivo* human trophoblast models, evidence also suggests that the placenta may independently regulate iron transfer to the fetus in specific circumstances.<sup>11</sup> A reduction of ferroportin expression on the apical fetal-facing membrane of placental syncytiotrophoblasts and increased expression of transferrin receptor 1 (TFR1) on the maternal-facing side have been observed in the setting of maternal iron deficiency. Placental metabolic homeostasis is subsequently maintained, protecting the fetus from the more detrimental condition of placental dysfunction.<sup>11</sup>

In laboring mothers at term, cord levels of IL-6 increase 4-fold even in the absence of evident infection.<sup>12</sup> Since the placenta is impermeable to IL-6,<sup>12,13</sup> these high cord blood levels suggest that labour could be associated with a fetal-neonatal inflammatory response, potentially triggered by labour-related mechanisms or exposure to infectious agents. Immediately after delivery, newborns face the most complex multi-organ physiological adaption that they will experience in their entire lives. Increased IL-6 levels in the newborn are thought to assist with organ system transition at birth (e.g. cytokine-induced synthesis of lung surfactant proteins<sup>14</sup>) and the activation of the immune system in the newborn<sup>15</sup>. IL-6 is also known to activate the JAK-STAT pathway, leading to the induction of hepcidin synthesis.<sup>16</sup> In previous studies we<sup>17,18</sup> and others<sup>19–22</sup> have demonstrated a rapid and profound hypoferremia occurring within the first few hours after delivery. This is assumed to have evolved as a defence against early-onset neonatal sepsis (EONS) and remains robust in premature and low-birthweight babies.<sup>18</sup> Several studies have shown that post-natal peripheral hepcidin and prohepcidin (precursor) levels are higher than those in cord blood.<sup>9,19,20</sup> We previously observed an initial increase in hepcidin within the first 12 hours of life in healthy newborns, positively correlated to raised IL-6 levels,<sup>17</sup> and we confirmed this in low birth weight and premature newborns.<sup>18</sup> Our data suggested that the IL-6-hepcidin-ferroportin axis plays a partial, but probably not exclusive, role in orchestrating the hypoferremia.

Here, we examine the duration of the hypoferremia and the likely regulatory influences in full-term, normal birthweight babies by analysing serial blood samples over the first week of life.

## 6.3 SUBJECTS AND METHODS

Full details of the Neolnnate Study ([clinicaltrials.gov](https://clinicaltrials.gov), NCT03353051) can be found in the published protocol paper.<sup>23</sup>

### Study Design

The Neolnnate Study tested whether preterm and/or low birthweight babies were capable of inducing acute hypoferremia previously noted in full-term, normal birthweight babies. Results of the primary outcomes have been presented elsewhere.<sup>18</sup> Here we describe the pre-planned secondary analysis of longitudinal changes in iron, hematological and inflammatory parameters over the first week of life within the term, normal weight babies from the FTB/NBW group. All babies were sampled from the cord artery (CDA) and vein (CDV) and had an early postnatal draw (V1) at 6-24h. For the longitudinal analysis over the first week of life and to avoid more than two blood draws per baby, the babies were then randomly allocated to a second blood draw at 25-80h (V2), 81-136h (V3) or 137-192h (V4) (**Figure 1**). Randomisation of second blood draw group allocation was completed using a random number calculator (GraphPad QuickCalcs, GraphPad Software INC, CA 92037, USA). This allocation was adapted according to study working hours and access to the newborn. Data collection started on the 5<sup>th</sup> July 2017 and ended on 1<sup>st</sup> February 2019.

### Ethics, standards and informed consent

The trial was approved by the Medical Research Council Unit The Gambia at London School of Hygiene and Tropical Medicine (MRCG at LSHTM) Scientific Coordinating Committee, the Joint Gambia Government/MRC Ethics Committee (SCC1525) and the London School of Hygiene and Tropical Medicine Ethics Committee (Ref:14316) and conducted according to Good Clinical Practice (GCP) standards. All participants gave written, informed consent.

## **Study setting**

Study participants were recruited from Kanifing General Hospital (formerly Serrekunda General Hospital), in the urban Kanifing region of The Gambia, West Africa.

## **Recruitment, screening and enrolment**

We planned to enrol 300 neonates into this longitudinal arm of the Neolnnate Study. For inclusion in this aim of the study, neonates were healthy, medically stable (i.e. not requiring resuscitation and with no signs of sepsis) with a gestational age  $\geq 37$  completed weeks (assessed by New Ballard Score<sup>24</sup>) and weighed  $\geq 2500$ g.

Pregnant mothers were excluded from the study if they were below the age of 18 years, had no fetal heartbeat detected upon admission, were known to be HIV-positive, were in receipt of TB therapy, had taken antibiotics in the last seven days, had a blood transfusion in the last month, were suffering from severe pre-eclampsia or antepartum haemorrhage, or were in another research study.

Babies were excluded at the delivery stage for the following reasons: major congenital malformations (not including polydactylism), blood transfusions given to mother or neonate, severe birth asphyxia (requiring resuscitation), neonates born via breech, vacuum or via caesarean section.

After the delivery stage, babies were excluded following the detection of infection or illness (i.e. information gained from a venous bleed or review of systems). Neonates were also removed from the study protocol if any medication other than intramuscular vitamin K, tetracycline eye ointment or immunisations was given. All medications given to mothers and neonates during the study period were recorded. Mothers who delivered multiple newborns were invited to enrol one of their neonates into the study.

## **Sample collection**

Once the neonate was fully delivered, one-minute delayed cord clamping was used (following World Health Organisation (WHO) policy<sup>25</sup>). After the umbilical cord was removed and cleaned, a trained study nurse identified the umbilical arteries (CDA) and umbilical vein (CDV). Blood was collected from each using separate blood draw equipment.

At 6-24 hours post-delivery, recruited mothers and their neonates were invited to a private consultation with the study research clinician. Demographic data were collected, along with a complete review of systems of the mother and neonate, and newborn anthropometry. Neuromuscular and physical maturation of each neonate was assessed using the New Ballard Score.<sup>24</sup> Immediately after passing the health assessment, a 3.5ml venous blood draw was performed on all neonates (V1).

During the community visit at the home of the neonate, a review of systems in the mother and child were conducted by a research nurse. This was followed by collecting data on medication, behaviour and immunisations of the neonate after leaving the hospital. A further sample of 3.5ml venous blood was then collected (V2-4) if the neonate was regarded as healthy.

## **Laboratory analyses**

A full haematology panel (using a Medonic M20M GP, Boule Diagnostics, Spanga, Sweden) and glucose-6-phosphate dehydrogenase deficiency test (R&D Diagnostics Limited, Papagos, Greece) were conducted on fresh whole blood. Serum was separated and stored at -20°C prior to analysis of ferritin, iron, unsaturated iron-binding capacity (UIBC), soluble transferrin receptor (sTfR), transferrin, c-reactive protein (CRP), haptoglobin, and alpha-1-acid glycoprotein (AGP) using a fully automated biochemistry analyser (Cobas Integra 400 plus, Roche Diagnostic, Switzerland). Transferrin saturation (TSAT) was calculated. Serum

samples were assessed for hepcidin concentration by ELISA (hepcidin-25 (human) EIA Kit, DRG, USA) with a dynamic range of 0.135-81 ng/mL.

In order to ensure a consistent assessment of haemolysis in all serum samples, batches of samples were thawed before entering the biochemistry analyser and visually scored by a single operator. A previously published specimen integrity chart for haemolysis was used as reference.<sup>26</sup> Samples were scored 0 (yellow 0 g/L of hemoglobin) to 6 (dark red 8 g/L of hemoglobin). Samples scoring  $\geq 5$  were removed from the analysis.

### **Sample size determination**

Sample size calculations for the primary outcomes of the Neolnate Study were based on data from a previous study<sup>17</sup> and are summarised in the protocol paper.<sup>23</sup> The secondary outcomes presented here were not subjected to a formal sample size analysis.

### **Statistical analysis**

Statistical analysis and preparation of figures were conducted using STATA v15.1 (Stat-Corp LP, College Station, TX, USA), DataDesk version 7.0.2 (Data Description Inc) and GraphPad Prism (GraphPad Software INC, CA 92037, USA). For continuous variables, baseline characteristics are presented as means ( $\pm$  SD) for normally distributed variables. All skewed data (hepcidin, CRP, AGP, sTfR and ferritin) were transformed using the ladder command in STATA. The ladder command searches a subset of the ladder of powers for a transformation that converts the variable of interest into a normally distributed variable. Results were confirmed graphically by the gladder command. Categorical variables are reported as proportions (%). Iron and inflammation markers were compared using ordinary least squares analysis of variance (OLS ANOVA), using Scheffé's post-hoc tests to control for multiple testing (V1 vs V2, V3 or V4). All hypothesis tests were two-sided at a significance level of 0.05. The rate of missing data was small (<5%), thus we did not impute missing data. Weighted Pearson network analysis was conducted using the Shiny Network



Application (<https://jolandakos.shinyapps.io/NetworkApp/>). The network was formatted using the Fruchterman-Reingold Algorithm<sup>27</sup> only showing moderate or strong (>0.3) associations between nodes. V2, V3 and V4 nodes were combined to aid visualisation.

## 6.4 RESULTS

A CONSORT diagram summarising subject recruitment is shown in **Figure 2**. There were 278 neonates with paired cord blood and V1 samples. Of these, 224 provided a second venous blood sample (V2, V3 or V4).

### Neonatal characteristics

Baseline characteristics are shown in **Table 1**. Newborns were healthy vaginally delivered babies, with a mean gestational age of  $39.4 \pm 1.3$ wk and mean birthweight of  $3299 \pm 368$ g. Many of the mothers (81.7%) received iron and folic acid during pregnancy as per WHO guidelines. Mean anthropometric measurements of all neonates fell within the 25th and 75th centiles of the WHO growth charts for gestational age.<sup>28</sup> The mean times of bleed were: V1 =  $12.7 \pm 0.32$ h; V2 =  $57.6 \pm 15.6$ h; V3 =  $105.8 \pm 17.0$ h; and V4 =  $156.7 \pm 21.0$ h.

### Changes in iron and chaperone proteins in the first week of life

Iron metabolism parameters over the first week of life can be seen in **Table 2** and **Figure 3**. Following the acute hypoferrremia on Day 1 (V1 =  $7.3 \pm 4.6$ µmol/L vs CDV =  $22.7 \pm 7.0$ µmol/L,  $P < 0.0001$ ) there was a steady increase in serum iron to  $16.5 \pm 3.9$ µmol/L at V4 ( $P$  for trend  $< 0.0001$ ) (**Fig 3A**). There was a slight decrease in transferrin levels (**Fig 3B**) but TIBC levels remained relatively constant (**Fig 3C**) and hence TSAT levels mirrored those for serum iron (**Fig 3D**). Following the acute postnatal hypoferrremia (V1 =  $14.4 \pm 6.0\%$  vs CDV =  $50.2 \pm 16.7\%$ ,  $P < 0.0001$ ) TSAT increased steadily to  $36.7 \pm 9.2\%$  at V4 ( $P$  for trend  $< 0.0001$ ). UIBC consequently decreased from V1 to V4 (from  $44.1 \pm 18.0$ µmol/L to  $29.5 \pm$

9.6 $\mu$ mol/L, P for trend <0.0001) (**Fig 3E**). Ferritin levels were very high at V1 (393  $\pm$  313 $\mu$ g/L) and showed a non-significant decline to 355  $\pm$  182 $\mu$ g/L at V4 (**Fig 3F**).

### **Changes in hepcidin and inflammatory markers in the first week of life**

Hepcidin levels in cord blood were high (19.4  $\pm$  14.4ng/ml) and doubled immediately after birth to reach 38.9  $\pm$  23.9ng/ml at V1 (P<0.0001). There was then a slight non-significant dip followed by a steady rise to 45.2  $\pm$  19.1ng/ml at V4 (P for trend <0.0001) (**Fig 4A**). CRP levels were low in cord blood (0.2  $\pm$  0.7 mg/L) and increased 10-fold immediately after birth (to 2.3  $\pm$  4.1mg/L at V1, P<0.0001) followed by a further steep rise to 5.6  $\pm$  9.3mg/L at V2 (P<0.0001 vs V1) and then a decline to 0.9  $\pm$  1.5 by V4 (P<0.0001 for trend)(**Fig 4B**). AGP showed a small increase after birth and then a further slow rise to V4 (P<0.0001) (**Fig 4C**). The heme-binding inflammatory-response protein haptoglobin increased between CDV and V1 (from 0.02  $\pm$  0.1g/L to 0.1  $\pm$  0.2g/L, P<0.0001) and then declined slightly to 0.08  $\pm$  0.1g/L at V4 (P for trend <0.0001) (**Fig 4D**).

### **Changes in hematological indices in the first week of life**

Cord hemoglobin (15.1  $\pm$  2.3g/dl) and hematocrit (42.2  $\pm$  7.0%) were high and there was a further increase by V1 (hemoglobin to 19.5  $\pm$  2.9g/dl, hematocrit to 55.0  $\pm$  9.7%, P<0.0001 for both increases). This was followed by a steady decline back towards cord-like levels by V4 (hemoglobin to 16.9  $\pm$  2.9g/dl, hematocrit to 46.9  $\pm$  9.0%, P<0.0001 for trend in both cases) (**Fig 5A,5B**). Soluble transferrin receptor levels dropped markedly reflecting the fact that there was a net breakdown of haemoglobin during the first week of life and an abundance of circulating iron to supply the needs of any on-going erythropoiesis (**Fig 5C**).

### **Pearson correlation network analysis of iron and inflammation markers**

Weighted Pearson correlation network analysis between the iron and inflammation markers in all samples (CDV, V1 and V2-4) are shown in **Figure 6**. The unconnected nodes for birthweight and gestational age reflect their lack of influence on any of the markers. As

expected, there were consistent associations between serum iron, TSAT, UIBC and TIBC at all time points, but the V1 nodes were notably separated from CDV and V2-V4 underscoring the unusual nature of the immediate post-natal hypoferremia. Hemoglobin and hematocrit followed a similar pattern, with cord values only correlating to V2-V4 values. The inflammatory markers CRP, AGP and haptoglobin were generally associated as would be expected (see lower part of the network) but ferritin, also an inflammatory marker, was notably separate. The most surprising feature was that hepcidin featured on the periphery of the network and was not associated with the inflammatory markers or serum iron.

### **Comparisons of iron and inflammation markers in arterial (CDA) and venous cord blood (CDV)**

**Figure 7** illustrates the comparisons between iron and inflammation markers in venous and arterial cord blood. For most markers there were no differences. TSAT was slightly higher in venous blood ( $50.5 \pm 1.0\%$ ) than arterial cord blood ( $46.9 \pm 1.0\%$ ,  $P < 0.0001$ ). Conversely, arterial blood had higher ferritin ( $276.9 \pm 14.7$  vs  $215.7 \pm 10.1 \mu\text{g/L}$ ,  $P < 0.0001$ ), UIBC ( $27.1 \pm 0.7$  vs  $23.4 \pm 0.7 \mu\text{mol/L}$ ,  $P < 0.0001$ ) and TIBC ( $49.8 \pm 0.57$  vs  $46.1 \pm 0.51 \mu\text{mol/L}$ ,  $P < 0.0001$ ). There were no differences in any red blood cell indices between venous and arterial cord blood, but white cell counts (with the exception of granulocyte count) were significantly higher in arterial blood (**Table 3**). Conversely platelet counts were significantly higher in venous blood.

## **6.5 DISCUSSION**

This study demonstrates that, following the previously reported<sup>17,18</sup> acute postnatal hypoferremia, serum iron and TSAT steadily revert towards cord-like levels over the first week of life. We interpret this as further evidence that the early hypoferremia has evolved to

help newborns navigate the hazardous journey as they progress from an almost sterile intra-uterine existence to the heavy bacterial exposures of extra-uterine life.

Our prior analyses<sup>17,18</sup> using data from independent studies, revealed that the early hypoferremia was, at least in part, driven by an inflammatory response to the birth process, eliciting a rapid IL-6-mediated rise in hepcidin. Hepcidin blocks the release of iron from enterocytes and macrophages<sup>1</sup> and thereby reduces serum iron through the dual actions of preventing iron absorption and recirculation. In neonates who receive insignificant amounts of dietary iron on Day 1, the latter mechanism dominates and the hypoferremia represents a temporary redistribution of iron away from the extracellular plasma where it would enhance the growth of any ingressing bacteria or fungi.<sup>17</sup> It is likely that additional hepcidin-independent mechanisms play an additional role in the hypoferremia.<sup>29</sup>

A surprising element of the current data is that iron and TSAT levels start to revert to normal despite hepcidin levels that continue to rise over the first week reaching values that are 3-4-fold higher than observed in healthy adults.<sup>30</sup> Furthermore, there was no correlation between hepcidin and serum iron or TSAT in the V2-4 samples. This unexpected disconnect between the relatively high levels of serum iron and TSAT coupled with high hepcidin concentrations suggests that early neonatal iron metabolism is desensitised to the action of hepcidin and thus the sequestration of intracellular iron is not maintained. This could be because the intracellular iron pools are saturated in the early post-partum period. We propose that macrophage cellular iron pools are increased in the first hours of life, initially due to the physiological hemolysis of fetal erythrocytes,<sup>31</sup> followed by the uptake of transferrin-iron complexes.<sup>32</sup> Erythrophagocytosis and the recycling of iron released from the catabolism of heme also add to intracellular iron levels.<sup>33</sup> This is further exacerbated by the effects of inflammation-induced hepcidin excess at 6-24 hours post-delivery, leading to hepcidin-induced co-degradation<sup>34</sup> and/or hepcidin occlusion<sup>35</sup> of the transmembrane iron transporter, ferroportin. We propose the initial hepcidin levels reduce expression of ferroportin on

macrophage cell membranes, thus eliciting the immediate post-natal hypoferremia, but that complete removal of all ferroportin molecules from the cell membrane is not achieved. This is supported by *in vivo* hepcidin challenge experiments showing a halving of ferroportin expression within 4 hours.<sup>36</sup> However, the complete removal of ferroportin was not achieved. Previous authors have suggested that excess levels of hepcidin more likely result in the blocking of the central cavity of ferroportin, rather than its internalization.<sup>35</sup> We propose that as hepcidin levels are consistently high during the first week of life, intermittent binding and releasing of hepcidin to the central cavity of individual ferroportin molecules may allow for slow rates of iron efflux into the circulation. We hypothesize that this results in a gradual increase in serum iron and TSAT, even in the high hepcidin environment, as seen in **Figures 3 and 4**. Previous research suggests that a potential cause of occlusion over internalization and the differences between tissues in sensitivity to hepcidin could be due to the differing endocytic machinery present in different cell types, or differing ferroportin glycosylation.<sup>36,37</sup> Further research in respect to neonatal macrophages is required.

Other studies have shown that following the neonatal period, circulating hepcidin levels decline to those similar to or lower than those observed in cord blood.<sup>38–40</sup> Increased expression of growth factors (i.e. IGF-1, HGF, EGF, PDGF-BB) is thought to be the cause of downregulation of hepcidin transcription.<sup>41,42</sup> Our research shows that this trend does not begin until after the first week of life.

Differences in hepcidin concentration between males and females are observed in adults<sup>43</sup> and children.<sup>44</sup> Previous research has noted significantly lower hepcidin concentrations in male neonates compared to females in the first week of life.<sup>45</sup> Though initially thought to be due to iron status and CRP concentration differences, it has recently been suggested that higher testosterone levels may be the cause. Though we did not measure testosterone levels in our samples, no significant difference in cord, V1 (6-24 hrs) or V2-4 (24-192 hours) hepcidin concentration was observed between sexes.

CRP levels peaked between 24-80 hours post-delivery, with again a surprising absence of correlation with hepcidin concentrations. This is despite the well-documented regulatory pathways of infection and inflammation on iron regulation.<sup>46</sup> Previous studies have suggested that the lack of correlation between hepcidin, IL-6 and CRP is due to differences in the kinetics of these proinflammatory biomarkers during the infection process. IL-6 concentrations spike very early in the course of infection or inflammation, hepcidin then follows and the rise of CRP is delayed and with a further delay to AGP release.<sup>47</sup>

The great majority of mothers in this study reported that they received iron and folic acid in pregnancy as per Gambian government guidelines but 52% remained anaemic in the last week before delivery. Despite this, ferritin levels in cord blood were high (CDV:  $213 \pm 158\mu\text{g/L}$ ) and levels almost doubled immediately after delivery to  $394 \pm 313\mu\text{g/L}$  at V1. It has previously been suggested that this is due to the physiological hemolysis of fetal red blood cells, which contain ferritin in high concentrations.<sup>22</sup> Similarly, we found elevated levels of haptoglobin, peaking at  $0.1 \pm 0.2\text{g/L}$  at 24-80 hours of life. We suggest this is another layer of nutritional immunity, as haptoglobin binds to hemoglobin, further restricting iron availability to invading microorganisms.<sup>48</sup>

We also recorded increases in newborn hemoglobin and hematocrit taken within the first hours of life (6-24 hours) compared to cord blood. This is likely to be in part due to postnatal dehydration, as well as vasomotor instability and venous pooling.<sup>49</sup> The decreases in hemoglobin and hematocrit over the first week of life are likely a response to the higher ambient oxygen concentration *ex-utero*.<sup>49</sup>

We undertook the comparative analysis of iron markers and inflammation in arterial and venous cord blood to ensure that non-standardized sampling of 'cord blood' in prior studies did not affect the comparisons between cord and postnatal samples. Serum iron was identical between sampling sites, and although there were significant differences for TIBC,

and hence reciprocally for TSAT, the differences were only in the order of 6-7%. Transferrin levels were similar, suggesting that differences in non-transferrin iron-binding compounds account for the small difference in TIBC and TSAT.

Cord arteriovenous samples show significant differences in all platelet parameters. This may be the result of the placenta being an active site of platelet production as previously suggested by Woods *et al.*<sup>50</sup>

There are several strengths and limitations to our study. The large sample size and the relative homogeneity of responses across most analytes provides confidence in the trends observed. A limitation is that maternal iron markers in mid-gestation, parturition and after delivery were not measured. This would have provided information as to what effect maternal iron and inflammation status had on the neonatal iron marker fluctuations we studied. Measurement of pro-inflammatory cytokines (e.g. IL-6 and IL-22) and growth factors (i.e. IGF-1, HGF, EGF and PDGF-BB) may have provided additional insights into the regulation of postnatal iron metabolism. Variables governing the hepcidin-independent regulation of iron redistribution are not yet known so could not be measured in our study. The effects of diurnal rhythm, iron supplementation and infection were not assessed and could be a direction for future research.

In conclusion, our results suggest that early postnatal hypoferremia is a fast-acting, yet short-lived adaptation likely to have evolved to protect the newborn from infection at the time they are most vulnerable. This is followed by a period of hepcidin desensitisation, or leaky intracellular iron sequestration, as iron efflux into the serum continues even in the presence of high serum hepcidin concentrations. The reduced need for iron for erythropoiesis during the first week of life could also increase the concentration of serum iron. This interpretation is supported by the observed decrease in sTfR levels over the first week indicating that erythroid tissues were not demanding iron. We have previously shown<sup>18</sup> that, in principle, the

duration of postnatal hypoferrremia might be extendable through the administration of mini-hepcidins as an ancillary tool against antimicrobial-resistant infections. The new data presented here suggests that any such intervention would need to overcome or circumvent the hepcidin resistance we report in the first week of life.

## 6.6 REFERENCES

- 1 Fisher AL, Nemeth E. Iron homeostasis during pregnancy. *Am J Clin Nutr* 2017; **106**: 1567S-1574S.
- 2 Koenig M, Tussing-Humphreys L, Day J, Cadwell B, Nemeth E. Hepcidin and Iron Homeostasis during Pregnancy. *Nutrients* 2014; **6**: 3062–83.
- 3 Awadallah SM, Abu-Elteen KH, Elkarmi AZ, *et al.* Maternal and Cord Blood Serum Levels of Zinc, Copper, and Iron in Healthy Pregnant Jordanian Women. *J Trace Elem Exp Med* 2004; **17**: 1–8.
- 4 Kocylowski R, Lewicka I, Grzesiak M, *et al.* Evaluation of Mineral Concentrations in Maternal Serum Before and After Birth and in Newborn Cord Blood Postpartum-Preliminary Study. *Biol Trace Elem Res* 2018; **182**: 217–23.
- 5 Lao TTT, Loong EPLP, Chin RKHK, Lam CWWK, Lam YMM. Relationship between newborn and maternal iron status and haematological indices. *Biol Neonate* 1991; **60**: 303–7.
- 6 Choi JW, Kim CS, Pai SH, J.W. C, C.S. K. Erythropoietic activity and soluble transferrin receptor level in neonates and maternal blood. *Acta Paediatr* 2000; **89**: 675–9.
- 7 Lipiński P, Styś A, Starzyński RR. Molecular insights into the regulation of iron metabolism during the prenatal and early postnatal periods. *Cell. Mol. Life Sci.* 2013; **70**: 23–38.
- 8 Rehu M, Punnonen K, Ostland V, *et al.* Maternal serum hepcidin is low at term and



- independent of cord blood iron status. *Eur J Haematol* 2010; **85**: 345–52.
- 9 Kulik-Rechberger B, Kościeszka A, Szponar EEE, *et al.* Hepcidin and iron status in pregnant women and full-term newborns in first days of life. *Ginekol Pol* 2016; **87**: 288–92.
  - 10 Lee S, Guillet R, Cooper EM, *et al.* Prevalence of anemia and associations between neonatal iron status, hepcidin and maternal iron status among neonates born to pregnant adolescents. *Pediatr Res* 2016; **79**: 42–8.
  - 11 Sangkhae V, Fisher AL, Wong S, *et al.* Effects of maternal iron status on placental and fetal iron homeostasis. *J Clin Invest* 2020; **130**: 625-640
  - 12 Duncombe G, Veldhuizen RAW, Gratton RJ, Han VKM, Richardson BS. IL-6 and TNF $\alpha$  across the umbilical circulation in term pregnancies: Relationship with labour events. *Early Hum Dev* 2010; **86**: 113–7.
  - 13 Aaltonen R, Heikkinen T, Hakala K, Laine K, Alanen A. Transfer of Proinflammatory Cytokines Across Term Placenta. *Obstet Gynecol* 2005; **106**: 802–7.
  - 14 Moss TJM, Newnham JP, Willett KE, Kramer BW, Jobe AH, Ikegami M. Early gestational intra-amniotic endotoxin: Lung function, surfactant, and morphometry. *Am J Respir Crit Care Med* 2002; **165**: 805–11.
  - 15 Protonotariou E, Chrelias C, Kassanos D, Kapsambeli H, Trakakis E, Sarandakou A. Immune response parameters during labor and early neonatal life. *In Vivo* 2010; **24**: 117–24.
  - 16 Wrighting DM, Andrews NC. Interleukin-6 induces hepcidin expression through STAT3. *Blood* 2006; **108**: 3204–9.
  - 17 Prentice S, Jallow AT, Sinjanka E, *et al.* Hepcidin mediates hypoferremia and reduces the growth potential of bacteria in the immediate post-natal period in human neonates. *Sci Rep* 2019; **9**: e16596
  - 18 Cross JH, Jarjou O, Mohammed NI, *et al.* Early postnatal hypoferremia in low birthweight and preterm babies. *EBioMedicine* 2019; **52**: 102613
  - 19 Balogh A, Szabo M, Kelen D, *et al.* Prohepcidin levels during human perinatal

- adaptation. *Pediatr Hematol Oncol* 2007; **24**: 361–8.
- 20 Kitajima J, Ohga S, Kinjo T, *et al.* Serum prohepcidin concentrations at birth and 1 month after birth in premature infants. *Pediatr Blood Cancer* 2011; **56**: 267–72.
  - 21 Yamada RRT, Leone CCR. Hematological and iron content evolution in exclusively breastfed late-preterm newborns. *Clinics* 2014; **69**: 792–8.
  - 22 Szabo M, Vasarhelyi B, Balla G, *et al.* Acute postnatal increase of extracellular antioxidant defence of neonates: the role of iron metabolism. *Acta Paediatr* 2001; **90**: 1167–70.
  - 23 Cross JH, Jarjou O, Mohammed NI, Prentice AM, Cerami C. Neonatal iron distribution and infection susceptibility in full term, preterm and low birthweight babies in urban Gambia: study protocol for an observational study. *Gates Open Res* 2019; **3**: 1469.
  - 24 Ballard JL, Khoury JC, Wedig K, Wang L, Eilers-Walsman BL, Lipp R. New Ballard Score, expanded to include extremely premature infants. *J Pediatr* 1991; **119**: 417–23.
  - 25 WHO | Guideline: Delayed umbilical cord clamping for improved maternal and infant health and nutrition outcomes. Geneva: World Health Organisation; 2014.  
<https://www.ncbi.nlm.nih.gov/books/NBK310511/>
  - 26 Dugan L, Leech L, Speroni KG, Corriher J. Factors Affecting Hemolysis Rates in Blood Samples Drawn From Newly Placed IV Sites in the Emergency Department. *J Emerg Nurs* 2005; **31**: 338–45.
  - 27 Fruchterman TMJ, Reingold EM. Graph drawing by force-directed placement. *Softw Pract Exp* 1991; **21**: 1129–64.
  - 28 Kiserud T, Piaggio G, Carroli G, *et al.* The World Health Organization Fetal Growth Charts: A Multinational Longitudinal Study of Ultrasound Biometric Measurements and Estimated Fetal Weight. *PLOS Med* 2017; **14**: e1002220.
  - 29 Enculescu M, Metzendorf C, Sparla R, *et al.* Modelling Systemic Iron Regulation during Dietary Iron Overload and Acute Inflammation: Role of Heparin-Independent Mechanisms. *PLoS Comput Biol* 2017; **13**: e1005322.

- 30 Diepeveen LE, Laarakkers CMM, Martos G, *et al.* Provisional standardization of hepcidin assays: creating a traceability chain with a primary reference material, candidate reference method and a commutable secondary reference material. *Clin Chem Lab Med* 2019; **57**: 864–72.
- 31 Buonocore G, Perrone S, Bracci R. Mechanisms involved in the increased hemolysis in the fetus and newborn. *Curr Pediatr Rev* 2017; **13**: 188-192
- 32 Sukhbaatar N, Weichhart T. Iron regulation: Macrophages in control. *Pharmaceuticals*. 2018; **11**: e137
- 33 Beaumont C, Canonne-Hergaux F. Erythrophagocytosis and recycling of heme iron in normal and pathological conditions; regulation by hepcidin. *Transfus. Clin. Biol.* 2005; **12**: 123–30.
- 34 Qiao B, Sugianto P, Fung E, *et al.* Hepcidin-induced endocytosis of ferroportin is dependent on ferroportin ubiquitination. *Cell Metab* 2012; **15**: 918–24.
- 35 Aschemeyer S, Qiao B, Stefanova D, *et al.* Structure-function analysis of ferroportin defines the binding site and an alternative mechanism of action of hepcidin. *Blood* 2018; **131**: 899–910.
- 36 Chaston T, Chung B, Mascarenhas M, *et al.* Evidence for differential effects of hepcidin in macrophages and intestinal epithelial cells. *Gut* 2008; **57**: 374–82.
- 37 Canonne-Hergaux F, Donovan A, Delaby C, Wang HJ, Gros P. Comparative studies of duodenal and macrophage ferroportin proteins. *Am J Physiol - Gastrointest Liver Physiol* 2006; **290**: G156-63.
- 38 Armitage AE, Agbla SC, Betts M, *et al.* Rapid growth is a dominant predictor of hepcidin suppression and declining ferritin in Gambian infants. *Haematologica* 2019; **104**:1542-1553
- 39 Stripeli F, Kapetanakis J, Gourgiotis D, Drakatos A, Tsolia M, Kossiva L. Post-transfusion changes in serum hepcidin and iron parameters in preterm infants. *Pediatr Int* 2018; **60**: 148–52.
- 40 Mupfudze TG, Stoltzfus RJ, Rukobo S, Moulton LH, Humphrey JH, Prendergast AJ.

- Hepcidin decreases over the first year of life in healthy African infants. *Br. J. Haematol.* 2014; **164**: 150–3.
- 41 Goodnough JB, Ramos E, Nemeth E, Ganz T. Inhibition of hepcidin transcription by growth factors. *Hepatology* 2012; **56**: 291–9.
  - 42 Sonnweber T, Nachbaur D, Schroll A, *et al.* Hypoxia induced downregulation of hepcidin is mediated by platelet derived growth factor BB. *Gut* 2014; **63**: 1951–9.
  - 43 Wisaksana R, de Mast Q, Alisjahbana B, *et al.* Inverse Relationship of Serum Hepcidin Levels with CD4 Cell Counts in HIV-Infected Patients Selected from an Indonesian Prospective Cohort Study. *PLoS One* 2013; **8**: e79904.
  - 44 Jaeggi T, Moretti D, Kvalsvig J, *et al.* Iron Status and Systemic Inflammation, but Not Gut Inflammation, Strongly Predict Gender-Specific Concentrations of Serum Hepcidin in Infants in Rural Kenya. *PLoS One* 2013; **8**: e57513.
  - 45 Uijterschout L, Domellöf M, Berglund SK, *et al.* Serum hepcidin in infants born after 32 to 37 wk of gestational age. *Pediatr Res* 2016; **79**: 608–13.
  - 46 Ganz T. Iron in innate immunity: starve the invaders. *Curr. Opin. Immunol.* 2009; **21**: 63–7.
  - 47 Lorenz L, Herbst J, Engel C, *et al.* Gestational Age-Specific Reference Ranges of Hepcidin in Cord Blood. *Neonatology* 2014; **106**: 133–9.
  - 48 Cassat JE, Skaar EP. Iron in infection and immunity. *Cell Host Microbe.* 2013; **13**: 509–19.
  - 49 Kates EH, Kates JS. Anemia and polycythemia in the newborn. *Pediatr. Rev.* 2007; **28**: 33–4.
  - 50 Woods MJ, Landon CR, Greaves M, Trowbridge EA. The placenta: A site of platelet production? *Platelets* 1992; **3**: 211–5.

**Abbreviations:**

AGP = Alpha 1-acid glycoprotein

CDA = Cord arterial blood

CDV = Cord venous blood

CRP = C-reactive protein

EDTA = Ethylenediaminetetraacetic acid

EGF = Epidermal growth factor

ELISA = Enzyme-linked immunosorbent assay

G6PD = Glucose-6-phosphate dehydrogenase

GCP = Good Clinical Practice

HGF = Hepatocyte growth factor

IGF-1 = Insulin-like growth factor-1

IL-6 = Interleukin 6

IL-22 = Interleukin 22

KGH = Kanifing General Hospital

KMC = Kanifing Municipal Council

MRCG = Medical Research Council Unit The Gambia at LSHTM

PDGF-BB = Platelet-derived growth factor - BB

ROS = Reactive oxygen species

SCC = Scientific Coordinating Committee

sTfR = Soluble transferrin receptor

TfR1 = Transferrin receptor

TIBC = Total iron-binding capacity

TSAT = Transferrin saturation

UIBC = Unbound iron-binding capacity

WHO = World Health Organisation

**Acknowledgements:**

We thank Kanifing Municipal Council (KMC) and Kanifing General Hospital (KGH) for their support during the study.

**Funding:**

The research was undertaken with a research grant provided by the Bill & Melinda Gates Foundation (OPP1152353). The funding agency had no role in the design and conduct of the study, nor in the collection, management, analyses or interpretation of the data or in the preparation, review, or approval of the manuscript. The Nutrition Theme of the MRC Unit The Gambia at LSHTM are supported by core funding MC-A760-5QX00 to the MRC Unit The Gambia/MRC International Nutrition Group by the UK MRC and the UK Department for the International Development (DFID) under the MRC/DFID Concordat agreement.

**Availability of Data and Materials:**

All data will be made available to researchers upon reasonable request to the study PI and clearance by the MRCG Scientific Coordinating and Ethics Committees.

**Authors' Contributions:**

**JHC** is a PhD student with MRCG at LSHTM. He contributed to the development of the protocols, study management, field data collection, development of the data analysis plan, conducting the data analysis and drafting of the manuscript.

**OJ** is a research clinician with MRCG at LSHTM. He contributed to the development of the protocols, study management, field data collection and patient care.

**NIM** is a statistician at MRCG at LSHTM. He wrote the data analysis plan in consultation with JHC, CC and AMP.

**SRG** was a visiting B.Sc. student from Cardiff University. He contributed to the laboratory analyses and field data collection.

**BJBT** is a laboratory technician with MRCG at LSHTM. He contributed to the laboratory analyses and field data collection.

**AMP** is a Professor of International Nutrition at LSHTM and Nutrition Theme Leader at MRCG at LSHTM. He conceived the study, obtained the funding, contributed to the development of the protocols, the data analysis plan, conducting the data analysis and the drafting of the manuscript.

**CC** is Senior Investigator Scientist at MRCG at LSHTM and Group Leader for Iron, Infection and Anemia. She was the PI on the trial, conceived the study, and obtained the funding. She was responsible for the overall development of the protocols, the data analysis plan, conducting the data analysis and the drafting of the manuscript.

All authors reviewed the final manuscript prior to submission.

#### **Competing Interests:**

The authors declare that they have no competing interests.

#### **Consent for Publication:**

Not applicable.

#### **Ethics Approval and Consent to Participate:**

This study has been approved by The Gambia Government/MRC Joint Ethics Committee (no. SCC1525) and London School of Hygiene and Tropical Medicine Ethics Committee (ref no. 14316). The study was conducted according to Good Clinical Practice (GCP) standards. The study procedures were explained to the neonate's mother/guardians orally and in writing. A neonate was only recruited into the study after the written, informed consent was provided by the mother/guardian.

**Author Details:**

<sup>1</sup>Medical Research Council Unit The Gambia at the London School of Hygiene & Tropical Medicine, Atlantic Boulevard, Fajara, P.O. Box 273, Banjul, The Gambia

<sup>2</sup>Cardiff University, Cardiff, Wales, CF10 3AT, United Kingdom



## Figure Legends:

**Figure 1: Study recruitment and blood draw design.** Mothers were approached on entering the KGH maternity ward, The Gambia. This was followed by the consenting process, recruitment and delivery data collection. At delivery, venous (CDV) and arterial (CDA) cord blood was collected after one-minute delayed cord clamping. The neonate was weighed after cord blood collection. At 6-24 hours post-delivery, the research study clinician conducted a health check of the mother and newborn. New Ballard Score was used to establish gestational age. If the neonate was deemed healthy, the V1 blood draw was completed. Follow-up in the community was conducted 24-216 hours post-delivery. This involved a health check of mother and newborn by the study nurse. If deemed healthy, the newborn was bled again (V2:  $\geq 24$  hours -  $< 80$  hours, V3:  $\geq 80$  hours -  $< 136$  hours, V4:  $\geq 136$  -  $< 192$  hours).

**Figure 2: CONSORT diagram for participant flow.** Two hundred and seventy-eight full-term, normal birthweight neonates were recruited to the study. Red outlined boxes represent all neonates excluded.

### **Figure 3: Changes in iron and chaperone proteins in the first week of life**

Blood was drawn from the umbilical cord vein (CDV) at birth and from the dorsum of the hand at serial time points for each individual ( $>6\text{-}\leq 24\text{hrs}$  and  $>24\text{-}\leq 216\text{hrs}$ ). Dots represent individual measurements. The blue line is a loess fit curve with 95% confidence intervals shaded in red. **A** = serum iron, **B** = transferrin, **C** = TIBC, **D** = TSAT, **E** = UIBC, **F** = ferritin.

### **Figure 4: Changes in hepcidin and inflammatory markers in the first week of life**

Blood was drawn from the umbilical cord vein at birth and from the dorsum of the hand at serial time points for each individual ( $>6\text{-}\leq 24\text{hrs}$  and  $>24\text{-}\leq 216\text{hrs}$ ). Dots represent individual

measurements. The blue line is a loess fit curve with 95% confidence intervals shaded in red. **A** = hepcidin, **B** = CRP, **C** = AGP, **D** = haptoglobin.

### **Figure 5: Changes in hematological indices in the first week of life**

Blood was drawn from the umbilical cord vein at birth and from the dorsum of the hand at serial time points for each individual (>6-≤24hrs and >24-≤216hrs). Dots represent individual measurements. The blue line is a loess fit curve with 95% confidence intervals shaded in red. **A** = hemoglobin, **B** = hematocrit, **C** = sTfR.

### **Figure 6: Weighted Pearson correlation network analysis of iron status and**

**inflammation variables between cord and post-natal blood samples.** Produced using the Shiny Network Application (<https://jolandakos.shinyapps.io/NetworkApp/>). This analysis is formatted into the “spring” layout that uses the Fruchterman-Reingold Algorithm <sup>27</sup> placing the more strongly correlated nodes closer together. Node colours define sample type (YELLOW = CDV, BLUE = V1, PINK = V2-4). The direction and size of the Pearson correlation between two nodes is represented using the colour (RED = Negative, GREEN = Positive) and thickness of an edge. Cord = umbilical cord blood. V1 = venous blood from dorsum of the hand at >6-≤24hrs. V2-4 = venous blood from dorsum of the hand at >24-≤216hrs. Fer = ferritin, Hep = hepcidin, Hct = hematocrit, Hgb = hemoglobin, GA = gestational age, BW = birthweight, CRP = C-reactive protein, Hapt = haptoglobin, AGP = alpha 1-acid glycoprotein, sTfR = soluble transferrin receptor, UIBC = unbound iron-binding capacity, TIBC = total iron-binding capacity, Trans = transferrin, fe = serum iron, TSAT = transferrin saturation.

**Figure 7: Comparison of iron status, inflammation and hematological parameters in cord arterial (CDA = BLUE) and venous (CDV = RED) blood.** Box plots represent the arithmetic mean with whiskers representing minimum and maximum values. \* =  $P < 0.05$ . No significance line =  $P > 0.05$ .

## Tables

**Table 1: Demographic, clinical and pregnancy outcome characteristics of the women and their newborns.**

Study Group	Exclusions	V1 (Total Sample)	V2	V3	V4
Number of Participants (n)	54	278	56	100	68
Characteristic	Mean (±SD)				
Gestational Age (Weeks)	39.4 (±1.3)	39.4 (±1.3)	39.4 (±1.4)	39.3 (±1.3)	39.6 (±1.2)
Birth Weight (g)	3361.2 (±377.9)	3299 (±368.3)	3270.4 (±355.3)	3293.1 (±364.7)	3281.9 (±378.7)
Head Circumference (cm)	34.5 (±1.0)	34.6 (±1.0)	34.5 (±1.0)	34.6 (±1.0)	34.6 (±1.2)
Length (cm)	50.0 (±1.6)	50.0 (±1.6)	50.2 (±1.5)	50.0 (±1.5)	49.9 (±1.7)
Maternal Hemoglobin ≤7 Days Before Delivery (g/dl)	11.6 (±2.2)	11.6 (±1.8)	11.4 (±1.8)	11.8 (±1.6)	11.6 (±1.8)
Age of Mother (Years)	27.5 (±6.8)	29.7 (±6.9)	31.2 (±7.3)	28.9 (±6.5)	31.4 (±6.8)
1 min APGAR Score (0-10)	9.5 (±0.8)	9.6 (±0.8)	9.6 (±0.7)	9.6 (±0.9)	9.6 (±0.7)
Time from Admission to Birth (Hours)	3.3 (±4.5)	3.2 (±5.2)	3.7 (±5.7)	2.7 (±4.1)	3.5 (±6.6)
Time from Delivery to V1 Blood Collection (Hours)	13.1 (±5.6)	12.7 (±5.4)	12.3 (±5.7)	13.1 (±5.0)	12.1 (±5.4)
Time from Delivery to V2 Blood Collection (Hours)	-	57.2 (±15.6)	57.6 (±15.6)	-	-
Time from Delivery to V3 Blood Collection (Hours)	-	105.8 (±17.0)	-	105.8 (±17.0)	-
Time from Delivery to V4 Blood Collection (Hours)	-	156.7 (±21.0)	-	-	156.7 (±21.0)
Percentage % (n)					
Percentage Male (%)	55.5% (30)	54.3% (151)	46.4% (26)	58.0% (58)	54.4% (37)
G6PD Deficiency Positive (%)	14.8% (8)	11.5% (32)	8.9% (5)	10.0% (10)	13.2% (9)
Multiple Births (%)	0.0% (0)	2.2% (6)	1.8% (1)	4.0% (4)	1.5% (1)
Percentage of Mother on Antenatal Iron/Folic Acid (%)	81.5% (44)	81.7% (227)	83.9% (47)	80.0% (80)	82.3% (56)

Data are presented as arithmetic mean (± SD) or as a proportion (%). Exclusions represent newborns successfully bled at cord and V1

timepoints but who were lost to follow up or sick before V2-4 blood draw was completed.

**Table 2: Comparison of iron status, inflammation and hematological parameters in cord and post-natal samples.**

Sample Type	n	CDV	n	V1	n	V2	n	V3	n	V4	V1 vs V2 (P value)	V1 vs V3 (P value)	V1 vs V4 (P value)	Trend (P Value)
Serum Iron (μmol/L)	275	22.7 (±7.0)	271	7.3 (±4.6)	54	10.1 (±3.2)	93	14.7 (±4.7)	62	16.5 (±3.9)	<b>0.0005</b>	<b>&lt;0.0001</b>	<b>&lt;0.0001</b>	<b>&lt;0.0001</b>
TSAT (%)	273	50.2 (±16.7)	271	14.4 (±6.1)	54	20.2 (±8.1)	91	32.2 (±13.0)	61	36.7 (±9.2)	<b>&lt;0.0001</b>	<b>&lt;0.0001</b>	<b>&lt;0.0001</b>	<b>&lt;0.0001</b>
Serum Hepcidin (ng/ml)	277	19.4 (±14.4)	270	38.9 (±23.9)	55	32.7 (±18.4)	92	45.9 (±17.7)	62	45.2 (±19.1)	ns	<b>0.003</b>	ns	<b>&lt;0.0001</b>
UIBC (μmol/L)	273	23.7 (±10.4)	271	44.1 (±18.0)	54	41.5 (±10.3)	91	32.4 (±10.2)	61	29.5 (±9.6)	ns	<b>&lt;0.0001</b>	<b>&lt;0.0001</b>	<b>&lt;0.0001</b>
TIBC (μmol/L)	273	46.3 (±8.1)	271	51.4 (±20.7)	54	51.6 (±9.8)	91	46.9 (±9.6)	61	46.0 (±10.5)	ns	ns	ns	<b>0.03</b>
Serum Ferritin (ug/L)	275	212.6 (±157.7)	266	393.9 (±312.6)	54	343.7 (±206.0)	93	311.8 (±131.3)	62	354.6 (±181.8)	ns	ns	ns	ns
Hemoglobin (g/dl)	270	15.1 (±2.3)	272	19.1 (±2.9)	49	19.5 (±2.9)	95	18.1 (±2.9)	64	16.9 (±2.9)	ns	<b>0.03</b>	<b>&lt;0.0001</b>	<b>&lt;0.0001</b>
Hematocrit (%)	270	42.2 (±7.0)	272	53.9 (±8.9)	49	55.0 (±9.7)	95	50.5 (±8.8)	64	46.9 (±9.0)	ns	<b>0.01</b>	<b>&lt;0.0001</b>	<b>&lt;0.0001</b>
Transferrin (g/L)	275	2.0 (±0.3)	273	2.0 (±0.3)	56	1.9 (±0.3)	93	1.7 (±0.2)	62	1.7 (±0.2)	ns	<b>&lt;0.0001</b>	<b>&lt;0.0001</b>	<b>&lt;0.0001</b>
Soluble Transferrin Receptor (mg/L)	273	6.0 (±2.0)	271	6.9 (±2.2)	54	6.6 (±2.0)	92	6.1 (±2.2)	62	5.1 (±1.3)	ns	<b>0.003</b>	<b>&lt;0.0001</b>	<b>&lt;0.0001</b>
Haptoglobin (g/L)	275	0.02 (±0.1)	273	0.03 (±0.07)	56	0.1 (±0.2)	93	0.08 (±0.2)	62	0.08 (±0.1)	<b>&lt;0.0001</b>	<b>0.001</b>	<b>0.01</b>	<b>&lt;0.0001</b>
Serum AGP (g/L)	275	0.2 (±0.1)	273	0.3 (±0.2)	56	0.4 (±0.2)	93	0.4 (±0.1)	62	0.4 (±0.1)	<b>&lt;0.0001</b>	<b>&lt;0.0001</b>	<b>&lt;0.0001</b>	<b>&lt;0.0001</b>
Serum CRP (mg/L)	275	0.2 (±0.7)	273	2.3 (±4.1)	55	5.6 (±9.3)	93	2.0 (±2.2)	62	0.9 (±1.5)	<b>&lt;0.0001</b>	ns	<b>0.01</b>	<b>&lt;0.0001</b>
Mean Corpuscular Volume (fl)	270	97.8 (±6.3)	272	97.7 (±6.0)	49	97.6 (±5.1)	95	96.0 (±5.9)	64	94.1 (±5.6)	ns	ns	<b>0.0002</b>	<b>&lt;0.0001</b>
Mean Corpuscular Hemoglobin (pg)	270	35.1 (±2.4)	272	34.8 (±2.4)	49	34.8 (±2.1)	95	34.5 (±2.2)	64	34.1 (±2.2)	ns	ns	ns	ns
Mean Corpuscular Hemoglobin Concentration (g/dl)	270	35.8 (±1.0)	272	35.6 (±1.1)	49	35.6 (±1.5)	95	35.9 (±0.9)	64	36.2 (±1.1)	ns	ns	<b>&lt;0.0001</b>	<b>0.0002</b>
White Blood Cell Count (unit/L)	269	13.3 (±5.7)	271	16.4 (±5.8)	48	10.5 (±5.4)	93	8.8 (±3.0)	64	9.2 (±2.5)	<b>&lt;0.0001</b>	<b>&lt;0.0001</b>	<b>&lt;0.0001</b>	<b>&lt;0.0001</b>
Lymphocyte Count (unit/L)	269	4.8 (±3.5)	271	4.2 (±2.9)	48	3.1 (±1.6)	93	3.3 (±1.2)	64	3.5 (±1.0)	<b>0.01</b>	<b>0.01</b>	ns	<b>0.0005</b>
Lymphocyte Percentage (%)	269	35.2 (±7.9)	271	26.5 (±8.8)	48	31.1 (±9.3)	93	38.7 (±7.6)	64	38.4 (±7.0)	<b>0.006</b>	<b>&lt;0.0001</b>	<b>&lt;0.0001</b>	<b>&lt;0.0001</b>
MID Cell Count (unit/L)	269	1.04 (±0.6)	271	1.54 (±1.0)	48	1.1 (±0.8)	93	1.2 (±0.8)	64	1.7 (±0.8)	<b>0.03</b>	<b>0.04</b>	ns	<b>0.0005</b>
MID Cell Percentage (%)	269	8.1 (±2.4)	271	9.8 (±5.6)	48	11.5 (±7.0)	93	14.8 (±7.8)	64	18.9 (±8.1)	ns	<b>&lt;0.0001</b>	<b>&lt;0.0001</b>	<b>&lt;0.0001</b>
Granulocyte Count (unit/L)	269	7.5 (±2.7)	271	10.7 (±4.2)	48	6.3 (±4.0)	93	4.2 (±1.9)	64	4.1 (±1.8)	<b>&lt;0.0001</b>	<b>&lt;0.0001</b>	<b>&lt;0.0001</b>	<b>&lt;0.0001</b>
Granulocyte Percentage (%)	269	56.6 (±8.7)	271	63.7 (±8.8)	48	57.4 (±9.3)	93	46.4 (±9.2)	64	42.7 (±9.1)	<b>0.0001</b>	<b>&lt;0.0001</b>	<b>&lt;0.0001</b>	<b>&lt;0.0001</b>
Red Blood Cell Count (unit/L)	270	4.3 (±0.7)	272	5.5 (±0.9)	49	5.6 (±1.0)	95	5.3 (±0.9)	64	5.0 (±0.9)	ns	ns	<b>0.0005</b>	<b>&lt;0.0001</b>
Red Blood Cell Distribution Width (%)	270	15.2 (±1.2)	272	15.4 (±1.4)	49	15.6 (±2.8)	95	15.1 (±0.9)	64	15.1 (±2.2)	ns	ns	ns	ns

Red Blood Cell Distribution Width - Absolute (fl)	270	80.7 (±8.9)	272	82.0 (±8.6)	49	82.5 (±15.7)	95	77.9 (±8.0)	64	75.3 (±7.0)	ns	<b>0.004</b>	<b>&lt;0.0001</b>	<b>&lt;0.0001</b>
Platelet Count (unit/L)	270	243.8 (±92.8)	272	258.7 (±94.0)	49	261.2 (±90.8)	95	234.8 (±97.0)	64	267.7 (±94.0)	ns	ns	ns	ns
Mean Platelet Volume (fl)	267	8.4 (±0.8)	271	8.5 (±0.8)	48	8.5 (±0.9)	93	8.7 (±0.8)	64	9.3 (±0.7)	ns	ns	<b>&lt;0.0001</b>	<b>&lt;0.0001</b>
Platelet Distribution Width (%)	267	43.5 (±3.0)	271	43.9 (±3.3)	48	44.5 (±3.5)	93	45.4 (±3.6)	64	46.4 (±2.7)	ns	<b>0.05</b>	<b>&lt;0.0001</b>	<b>&lt;0.0001</b>
Platelet Distribution Width - Absolute (fl)	267	11.8 (±1.3)	271	12.1 (±1.4)	48	12.1 (±1.6)	93	12.5 (±1.5)	64	13.3 (±1.2)	ns	<b>0.003</b>	<b>&lt;0.0001</b>	<b>&lt;0.0001</b>
Platelet Crit (%)	267	0.201 (±0.07)	271	0.215 (±0.07)	48	0.2 (±0.1)	93	0.2 (±0.07)	64	0.2 (±0.08)	ns	ns	ns	<b>0.009</b>
Plate Large Cell Ratio (%)	267	18.0 (±5.4)	271	19.2 (±5.8)	48	19.4 (±6.3)	93	21.1 (±5.9)	64	24.3 (±5.0)	ns	ns	<b>&lt;0.0001</b>	<b>&lt;0.0001</b>
Plate Large Cell Count (unit/L)	267	42.1 (±14.6)	271	46.9 (±17.0)	48	48.4 (±17.4)	93	47.6 (±17.8)	64	63.1 (±21.8)	ns	ns	<b>&lt;0.0001</b>	<b>&lt;0.0001</b>

Data are presented as mean (± SD) and analysed by ordinary least squares analysis of variance (OLS ANOVA). Number of available results differs by each parameter, due to limitations in blood sample volume for some participants. Scheffé's post-hoc tests were conducted between V1 vs V2, V3 and V3 groups. *P* values in bold font are considered significant based on *P*<0.05. Hepcidin (log), ferritin (log), sTfR (log), AGP (Sqrt) and CRP (log) were all transformed to form a normal distribution before ordinary least squares analysis of variance (OLS ANOVA) and reverse transformation for the listings above.

**Table 3: Comparison of iron status, inflammation and hematological parameters in cord venous and arterial blood.**

Sample Type	n	Cord (Arterial)	n	Cord (Venous)	<i>P value</i>
Serum Iron (μmol/L)	258	22.7 (±6.9)	275	22.7 (±7.0)	0.8
TSAT (%)	255	47.1 (±15.8)	273	50.2 (±16.7)	<b>&lt;0.0001</b>
Serum Hepcidin (ng/ml)	249	19.7 (±14.4)	277	19.4 (±14.4)	0.34
UIBC (μmol/L)	255	27.1 (±11.3)	273	23.7 (±10.4)	<b>&lt;0.0001</b>
TIBC (μmol/L)	257	49.8 (±9.1)	273	46.3 (±8.1)	<b>&lt;0.0001</b>
Serum Ferritin (ug/L)	241	277.0 (±234.2)	275	212.6 (±157.7)	<b>&lt;0.0001</b>
Hemoglobin (g/dl)	242	15.2 (±2.6)	270	15.1 (±2.3)	0.53
Hematocrit (%)	260	43.0 (±7.5)	270	42.2 (±7.0)	0.16
Transferrin (g/L)	258	2.0 (±0.3)	275	2.0 (±0.3)	0.48
Soluble Transferrin Receptor (mg/L) (log)	260	6.0 (±1.9)	273	6.0 (±2.0)	0.86
Haptoglobin (g/L)	260	0.02 (±0.05)	275	0.02 (±0.06)	0.64
Serum AGP (g/L)	260	0.2 (±0.1)	275	0.2 (±0.1)	0.97
Serum CRP (mg/L)	242	0.2 (±0.03)	275	0.2 (±0.7)	0.27
Mean Corpuscular Volume (fl)	241	98.2 (±6.2)	270	97.8 (±6.3)	0.07
Mean Corpuscular Hemoglobin (pg)	241	35.1 (±2.5)	270	35.1 (±2.4)	0.9
Mean Corpuscular Hemoglobin Concentration (g/dl)	241	35.8 (±1.1)	270	35.8 (±1.0)	0.09
White Blood Cell Count (unit/L)	241	14.2 (±6.1)	269	13.3 (±5.7)	<b>0.004</b>
Lymphocyte Count (unit/L)	241	5.5 (±3.7)	269	4.8 (±3.5)	<b>&lt;0.0001</b>
Lymphocyte Percentage (%)	241	38.0 (±9.1)	269	35.2 (±7.9)	<b>&lt;0.0001</b>
MID Cell Count (unit/L)	241	1.2 (±0.8)	269	1.0 (±0.6)	<b>&lt;0.0001</b>
MID Cell Percentage (%)	241	8.8 (±3.2)	269	8.1 (±2.4)	<b>0.0001</b>
Granulocyte Count (unit/L)	241	7.5 (±3.0)	269	7.5 (±2.7)	0.53
Granulocyte Percentage (%)	241	53.2 (±9.4)	269	56.6 (±8.7)	<b>&lt;0.0001</b>
Red Blood Cell Count (unit/L)	242	4.4 (±0.8)	270	4.3 (±0.7)	0.44
Red Blood Cell Distribution Width (%)	242	15.3 (±1.0)	270	15.2 (±1.2)	0.11
Red Blood Cell Distribution Width - Absolute (fl)	242	81.5 (±9.2)	270	80.7 (±8.9)	<b>0.008</b>
Platelet Count (unit/L)	242	193.0 (±86.7)	270	243.8 (±92.8)	<b>&lt;0.0001</b>
Mean Platelet Volume (fl)	237	8.6 (±0.8)	267	8.4 (±0.8)	<b>&lt;0.0001</b>
Platelet Distribution Width (%)	237	44.6 (±3.4)	267	43.5 (±3.0)	<b>&lt;0.0001</b>
Platelet Distribution Width - Absolute (fl)	237	12.3 (±1.4)	267	11.8 (±1.3)	<b>&lt;0.0001</b>
Platelet Crit (%)	237	0.2 (±0.07)	267	0.2 (±0.1)	<b>&lt;0.0001</b>
Plate Large Cell Ratio (%)	237	20.4 (±5.9)	267	18.0 (±5.4)	<b>&lt;0.0001</b>
Plate Large Cell Count (unit/L)	237	42.1 (±14.6)	267	42.1 (±14.6)	<b>&lt;0.0001</b>

Data are presented as mean (± SD) and analysed by one-way analysis of variance. *P* values

in bold font are considered significant based on *P*<0.05. Hepcidin (log), ferritin (log), sTfR

(log), AGP (Sqrt) and CRP (log) are all transformed to form a normal distribution before one-way analysis of variance (ANOVA) and reverse transformation for the listings above.

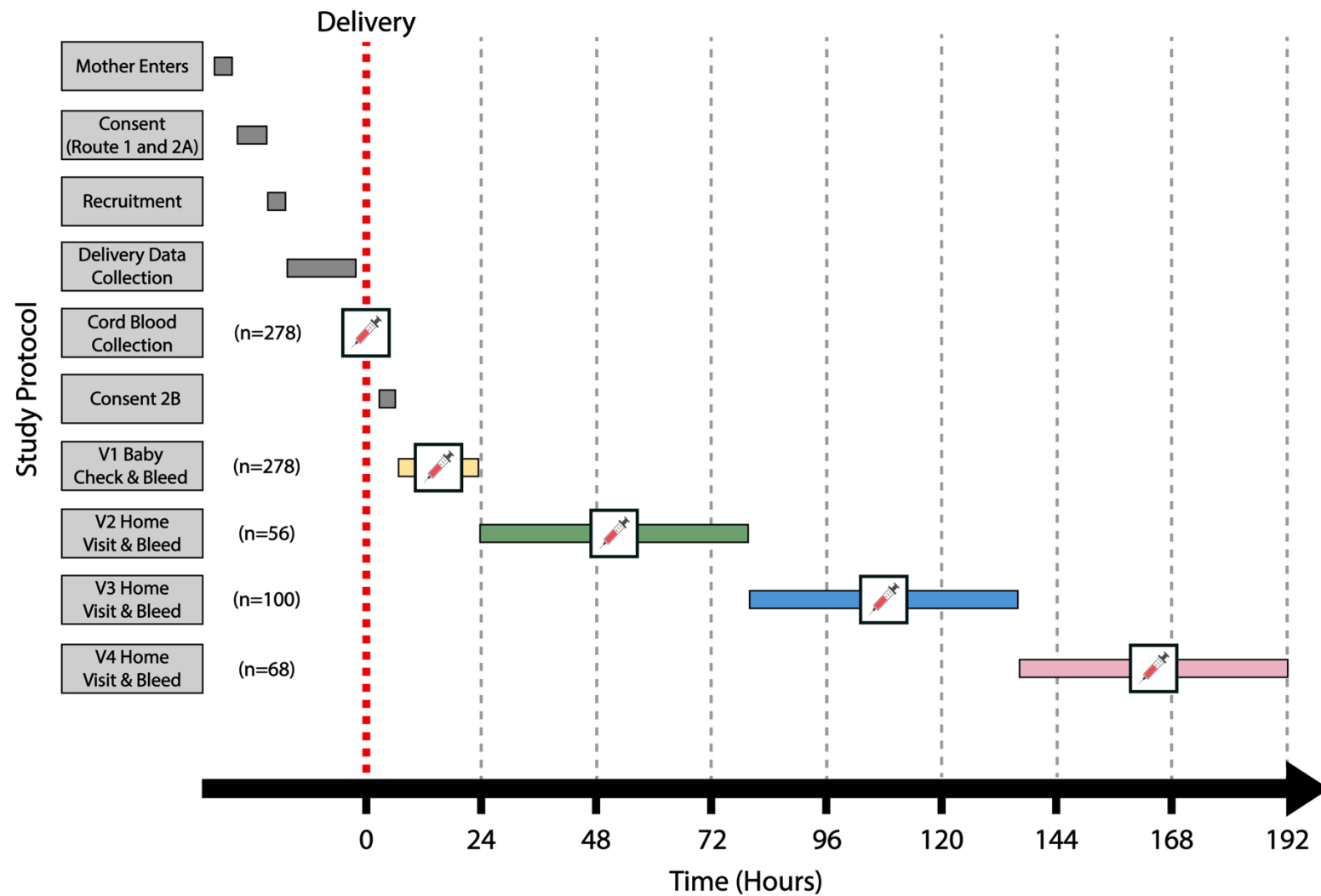
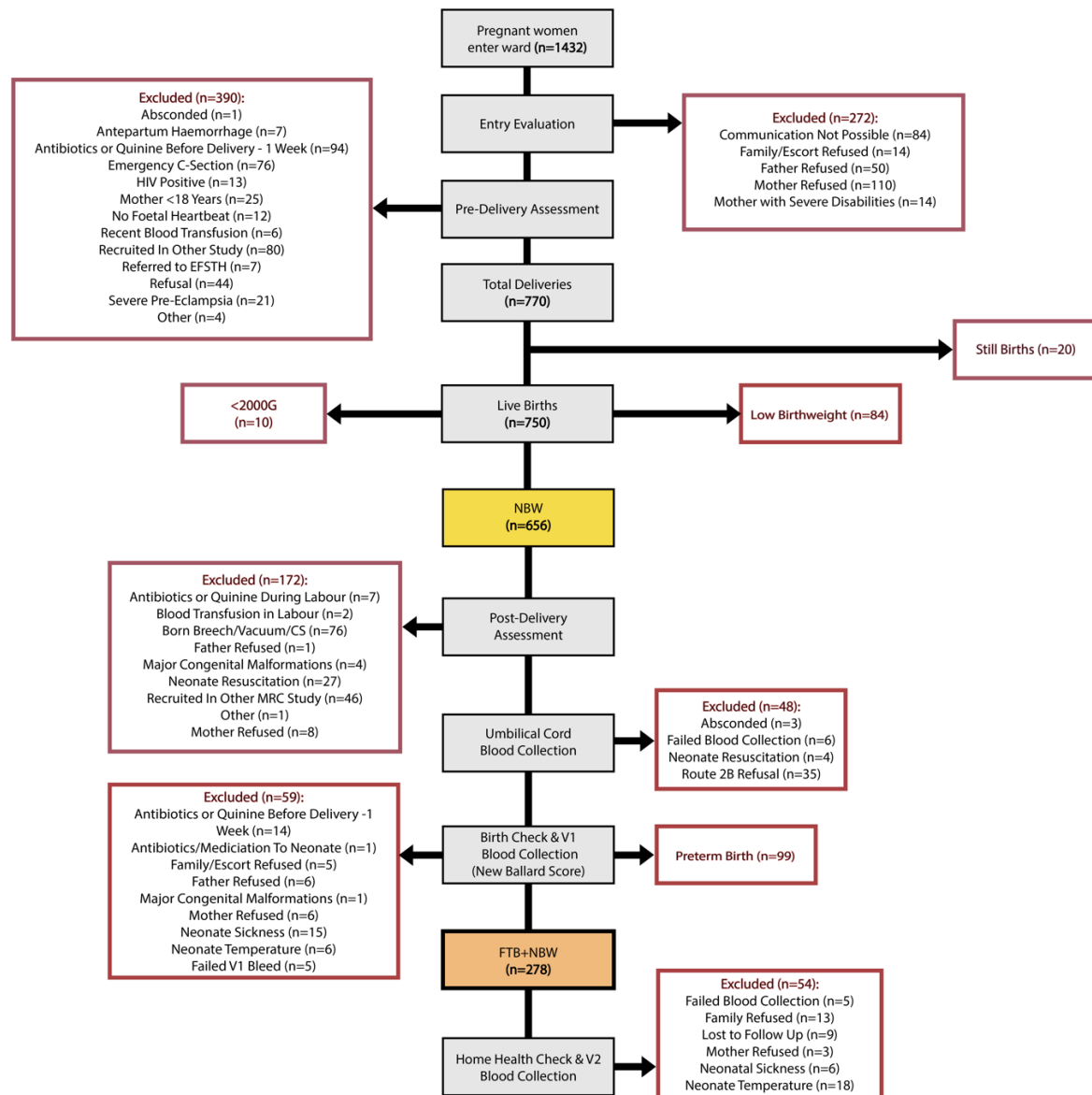
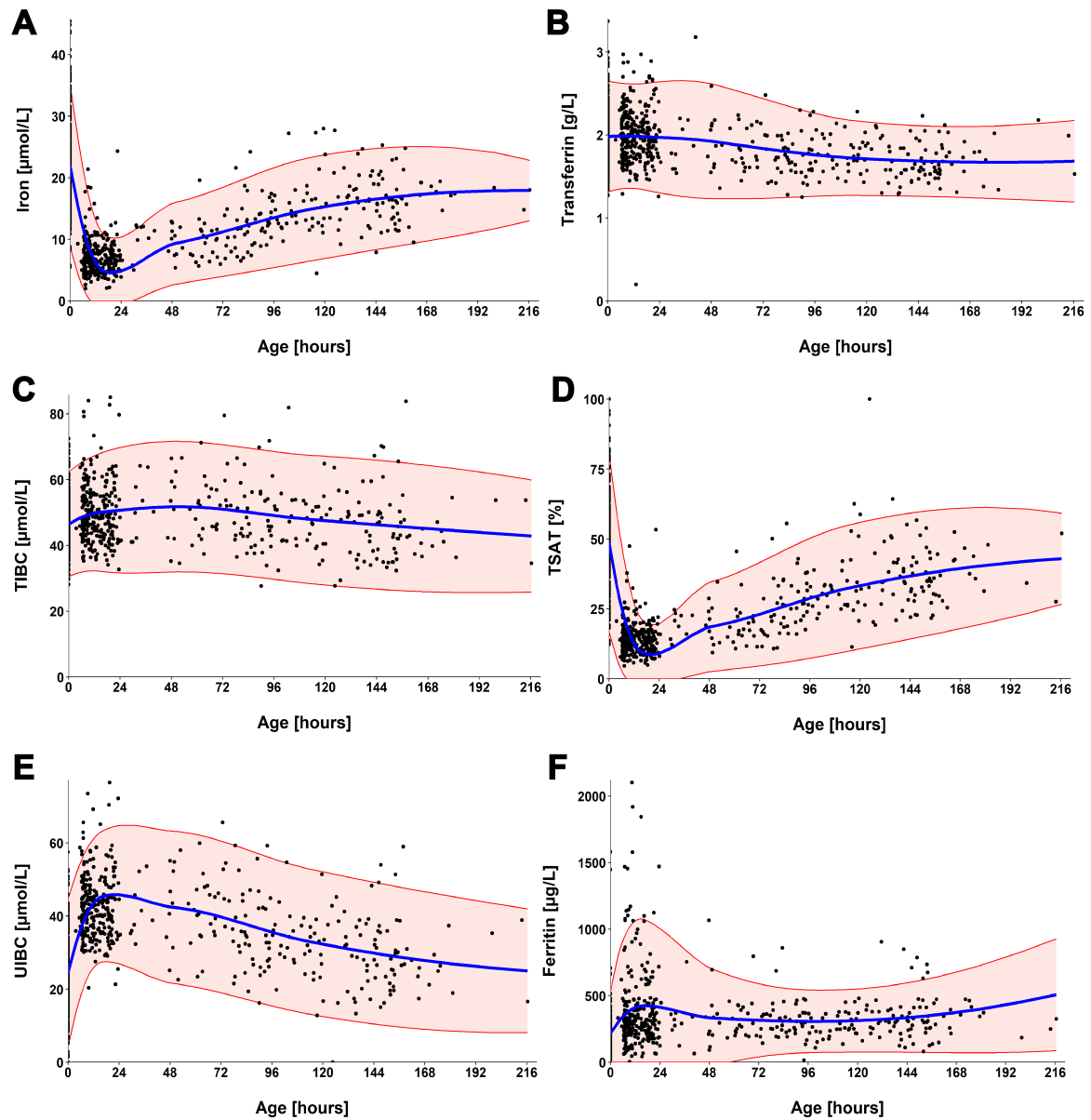


Figure 1: Study recruitment and blood draw design.

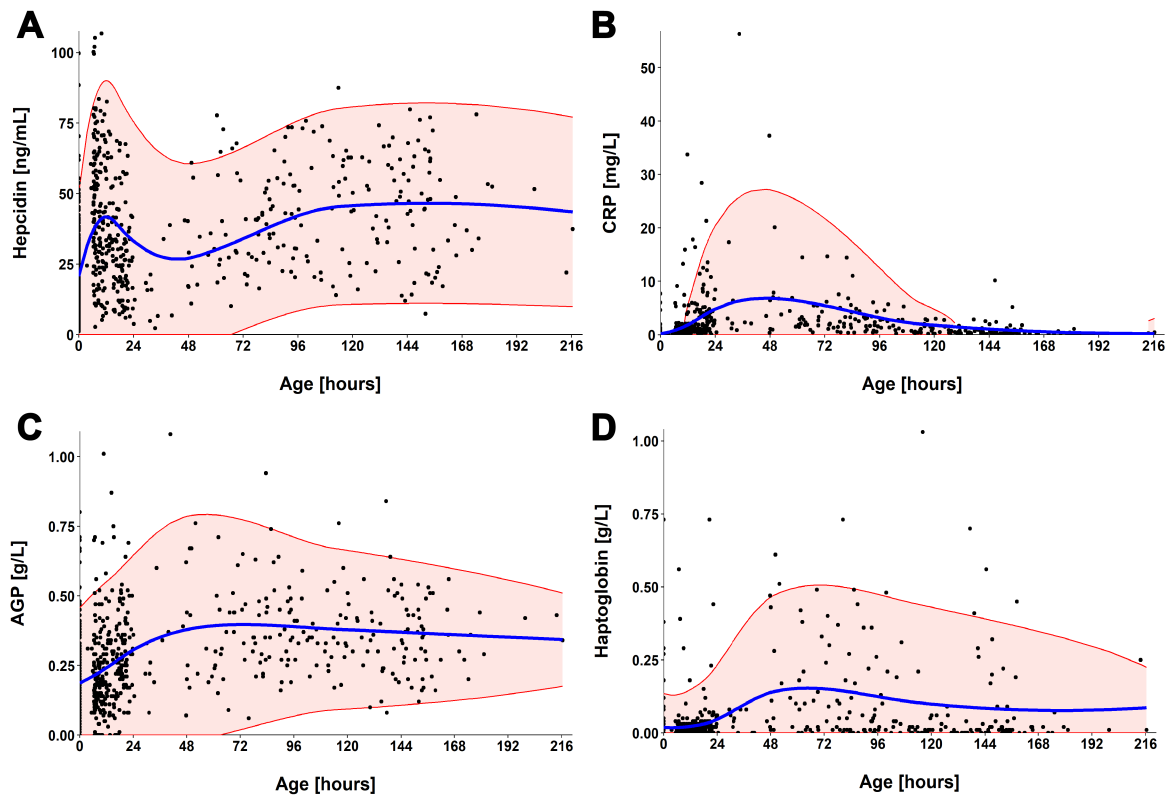




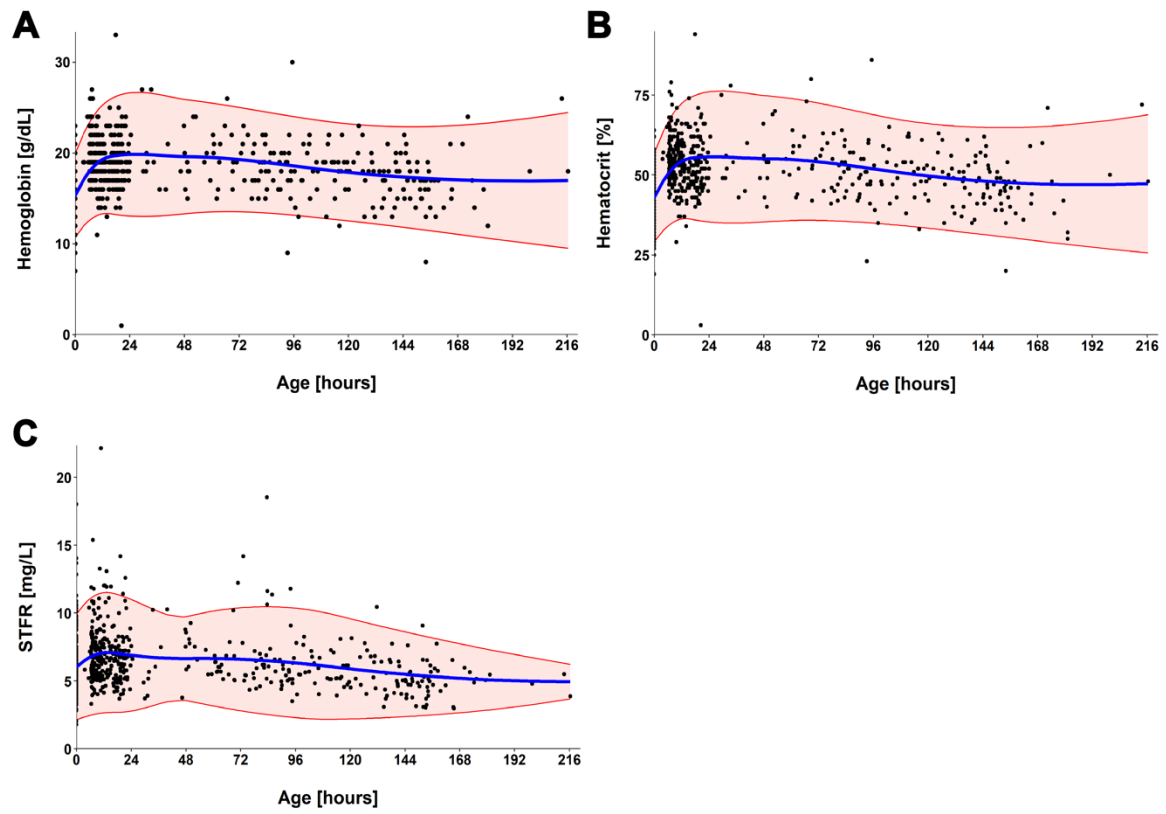
**Figure 2: CONSORT diagram for participant flow.**



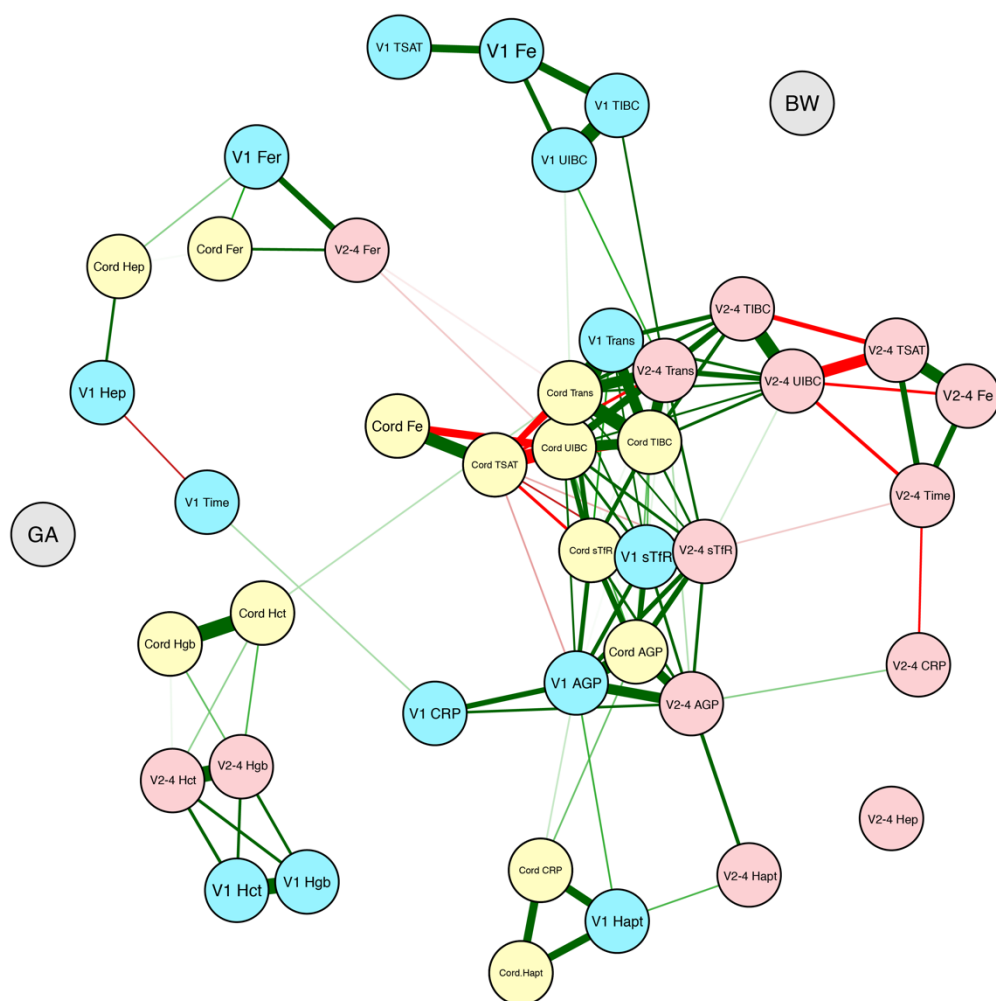
**Figure 3: Changes in iron and chaperone proteins in the first week of life.**



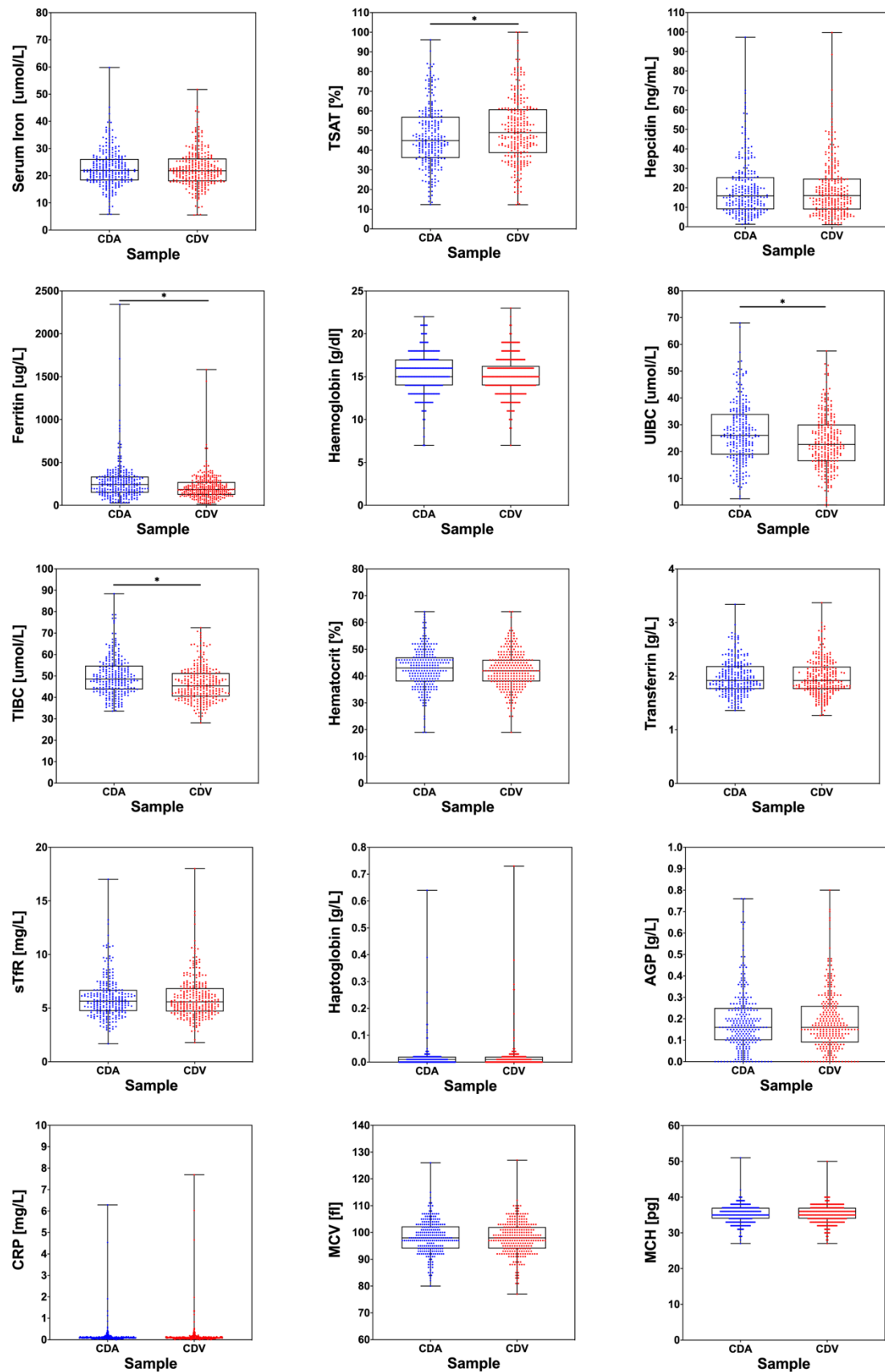
**Figure 4: Changes in hepcidin and inflammatory markers in the first week of life.**



**Figure 5: Changes in hematological indices in the first week of life.**



**Figure 6: Weighted Pearson correlation network analysis of iron status and inflammation variables between cord and postnatal blood samples.**



**Figure 7: Comparison of iron status, inflammation and hematological parameters in cord arterial (CDA = BLUE) and venous (CDV = RED) blood.**

## Chapter 7 - Discussion

---

### Summary of Chapter

This chapter focuses on describing a flowing narrative of the overall content of this thesis. It begins with a description of the essential aspects of study design, the critical learning gained from previous pilot studies and the literature review process. A detailed summary of what was discovered during the Neolnnate Study and how it addressed critical research gaps then follows.

The subsequent section then notes several of the main issues and learning points faced during the research study process. At each point in this section, we outline the problem, the science behind it and how our research team overcame it.

Finally, this discussion recommends four potential future research areas. This firstly focuses on the role the placenta plays in the maternal-fetal iron transfer process. This is followed by a review of possible triggers of the neonatal hypoferremic response (e.g. hepcidin-dependent and hepcidin-independent mechanisms), and what research is required in order to understand these mechanisms further. Thirdly, this section describes the current research into harnessing hypoferremia to combat other infections and how this area is moving forward. This includes the current development of therapies that use iron chaperone proteins (e.g. haptoglobin, lactoferrin and lipocalin-2), bacterial iron-uptake mechanisms, and hepcidin agonists to treat bacterial infections. Lastly, this chapter outlines how future research should focus on the role of macrophages in the neonatal hypoferremic response and the kinetics of hepcidin-resistance post-hypoferremia.

## 7.1 A Review of the Neolnnate Study

### 7.1.1 Neolnnate Study: Where Did It Begin? (Chapter 1)

In light of the evidence that suggests hepcidin-induced hypoferremia is protective against severe bacterial infections,<sup>1</sup> the findings of our team's pilot study were an exciting, unexpected finding. This PhD study was formulated after the previous research by Prentice *et al.* (HYPO-G study).<sup>2</sup> This study was a non-blinded randomised controlled trial conducted on 120 full-term, normal birthweight Gambian neonates. Infants were randomised to receive Bacillus Calmette–Guérin (BCG) vaccination on the day of birth or after study completion at four days of age. Blood samples for investigation of iron parameters, hepcidin, IL-6 and red blood cell parameters were collected at birth (i.e. umbilical cord) and time-points up to 96 hours of age. This study found that regardless of whether neonates received BCG vaccination or not, that during the first 24 hours of life full-term neonates actively reduce the serum iron concentration and transferrin saturation of the blood. This was found to occur as early as six hours postpartum and lasted for two to three days. This response was in strong correlation with hepcidin, suggesting that hepcidin regulates this response as seen in previous mechanistic studies.<sup>3</sup> This finding was also correlated to levels of IL-6, suggesting that inflammatory stimulation was the trigger. Using a micro-adaptation of the *ex vivo* bacterial growth assay previously described in our FeVir study,<sup>4</sup> we showed that the lack of iron similarly reduced growth rates of *E. coli*, *S. aureus*, *S. pneumoniae* and *S. agalactiae* (common neonatal sepsis causing pathogens) inoculated into venous samples collected at 6-24 hours of life, in comparison to cord samples. The growth rates were significantly associated with TSAT level in the serum.<sup>2</sup>

Supported by similar but smaller reductions in serum iron concentration and TSAT observed in the first week of life in other studies,<sup>5–8</sup> this observation suggested that the neonatal



hypoferremia may have evolved as a mechanism designed to protect neonates from infection during the first critical days of life. This is a result of limiting the pathogenicity and virulence of invading bacterial organisms. However, more diverse populations with a range of birthweights and gestational ages were required to confirm this. This is especially the case in the light of evidence that suggests preterm neonates suffer deficiencies in Toll-like receptors (TLRs), a vital component of the inflammation-induced hypoferremic response.<sup>9</sup> We further hypothesised that this might lead to a dampening of this response, or a complete failure to reduce systemic iron concentrations. We believed that this might, in part, explain why being premature and/or low birthweight may be risk factors for neonatal sepsis.<sup>10</sup>

### **7.1.2 A Review of Iron Homeostasis Over the First Month of Life (Chapter 3)**

This research studentship started with a systematic search of literature relating to hepcidin, serum iron and transferrin saturation levels in full-term and preterm newborns during the first month of life (**Chapter 3**). Retrieved data suggested that hepcidin, serum iron and TSAT levels for adults and infants are much lower than those observed in cord blood and venous blood during the first month of life. Accumulated data further strengthened our previously defined hypothesis that full-term neonates possess the ability to produce a hepcidin-mediated hypoferremic response post-delivery. Nonetheless, data concerning preterm neonates was lacking and subsequently it was unclear as to whether they produce a similar response. This dearth of studies is presumably due to their relatively unstable medical condition, and the complex ethical issues around bleeding premature neonates. This review also highlighted a lack of data from blood samples collected in the first hours of life (6-24 hours) and over the first week, even in term babies.

Previously, Van der Vorm *et al.* formulated a series of linear equations to assist in the standardisation of each commonly used hepcidin measurement assay.<sup>11</sup> In our review, we

used these equations to standardise results from previous publications. This allowed the calculation of weighted mean averages in cord and venous blood for full-term and preterm neonates. We believe the main strength of this review was that it is the first review to compare retrospective serum hepcidin concentrations between studies, using standardised values. This is the first review of hepcidin, serum iron and TSAT in cord blood and venous samples over the first weeks of life, that was stratified by gestational age. The learning points from this review for the following observational study were the importance of collecting of very early postnatal venous blood (6-24 hours), the use of the DRG hepcidin ELISA kit, the importance of detailed labour and delivery data, and standardisation of cord blood collection protocols (**Chapter 4**).

Weaknesses of the review included the allocation of each study group or population in each publication to a mean gestational age. This was due to limited access to raw study data. It is suspected that this has reduced any natural variation potentially caused by gestational age between the reviewed populations. Equally, due to studies rarely stating the average birth weight of each population, analysis stratified by birthweight was not possible. Furthermore, the retrieval of gestational age was a crucial aspect of the search strategy; however, few studies documented the method used to calculate it. Due to the substantial differences in the accuracy of different techniques,<sup>12</sup> the true mean gestational age of each population may vary by differing amounts.

### **7.1.3 Early Postnatal Hypoferremia in Low Birthweight and Preterm Babies (Chapter 5)**

Initiation of the Neolnnate Study then began, with the formation of a protocol and data analysis plan. The primary objective of the Neolnnate study was to ascertain if there was a defect in the ability of preterm and low birthweight newborns to elicit a hypoferremic

response immediately after delivery (**Chapter 5**). This observational study ended with the recruitment of 152 babies who were either preterm (PTB) (born  $\geq 32$  to  $< 37$  weeks gestational age) and/or low birth weight (LBW) ( $< 2500\text{g}$ ) (PTB/LBW) and 278 term (FTB), normal-weight babies (NBW) (FTB/NBW). Blood was sampled from the umbilical cord vein and artery, as well as matched venous blood samples taken from all neonates between 6-24 hours after delivery. In both PTB/LBW and FTB/NBW babies, we observed serum iron decreased 3-fold within 12 hrs of delivery compared to umbilical blood, and transferrin saturation showed a similar decline with a consequent increase in unsaturated iron-binding capacity. C-reactive protein levels increased over 10-fold and hepcidin levels doubled over the same period. We can now conclude that there was no difference in any of these responses between PTB/LBW and FTB/NBW babies. This result suggests that the innate postnatal iron restriction strategy in the first hours of life has evolved as an intrinsic mechanism to protect all neonates from common pathogens and/or free-radical damage, regardless of gestational age or birthweight. This finding now prompts further research into the use of iron restriction as a transient bacteriostatic mechanism to limit bacterial growth and virulence in other instances of infection.

One of the limitations of this study includes that we did not recruit mothers with complex medical histories (e.g. pre-eclampsia, antepartum haemorrhage or antenatal infection) or sick babies (e.g. with birth asphyxia or suspected sepsis). This was deliberate; however, the question remains as to whether neonates with suspected sepsis, have sepsis due (in part) to them lacking this innate immune response. This question will remain challenging to assess as it will be difficult to access the exact point of the initiation of the infection and to what degree any change in inflammation or iron restriction is due to the presence of the infection itself. Additionally, we did not recruit newborns born  $< 32$  completed weeks gestation and/or  $< 2000\text{g}$  birth weight, or those delivered via caesarean section, vacuum or forceps. As a result, we cannot speculate whether similar hypoferremic responses occur in these newborns. However, reciprocally, we can reason that the responses we observed in the

Neolnnate Study are a characteristic feature of normal human birth and not due to pathological circumstances.

#### **7.1.4 Iron Homeostasis in Full Term, Normal Birthweight Gambian Neonates Over The First Week of Life (Chapter 6)**

The final objective of the Neolnnate Study was to describe the longitudinal changes in iron metabolism and haematological parameters after the initial hypoferremic response at 6-24 hours post-delivery. This was achieved by the collection of a second venous sample from all full-term, normal birth weight newborns later in the first week of life. This resulted in 224 of the 278 full-term, normal birthweight neonates (FTB/NBW) providing a cord blood sample, and two peripheral venous blood samples at 6-24h and 24-192h of age. This publication (**Chapter 6**) highlights that levels of hepcidin quickly return to previous mean cord-like levels at 24-48 hrs, and then subsequently increase to very high levels at 80-136hrs post-delivery. Serum iron and TSAT levels also increased during this period. This observation suggests that the early postnatal hypoferremia is a fast-acting, yet short-lived, adaptation that is followed by a period of hepcidin desensitisation or leaky intracellular iron sequestration. We suggest that this is the explanation for increased iron efflux into the serum, even in the presence of high serum hepcidin concentrations. This increase in circulating serum iron could be heightened by the reduced need for iron for erythropoiesis over the same period. A marker of this is the decreasing sTfR level over the first week of life. Weaknesses in research directed toward this objective include the lack of data relating to the quality of breastfeeding during the first week of life. This would have allowed us to control the analysis of V2-4 samples for the effect of enteric iron absorption from breast milk or formula (though both are likely to contribute small amounts of iron).<sup>13</sup> Similarly, the measurement of maternal iron markers before (i.e. midgestation and parturition) and after delivery could have provided greater detail into the role of iron endowment between the mother and fetus and also the

part that labour-induced inflammation plays in the recorded hypoferremic response. Though maternal iron markers have been analysed in relation to cord iron markers,<sup>14,15</sup> we believe no analysis has been completed concerning the change in serum iron and TSAT we have recorded. This analysis could provide further evidence as to what is triggering this phenomenon. The measurement of pro-inflammatory cytokines (e.g. IL-6 and IL-22) and growth factors (e.g. IGF-1, HGF, EGF and PDGF-BB) in all samples would have provided greater insight into the regulation of neonatal iron homeostasis.

## **7.2 Neolnnate Study: Study Design, Issues Faced and Learning Points for the Future**

The Neolnnate study was a challenging study to design, due to its time-sensitivity, complicated exclusion criteria and complex ethical issues.

The review of the previous literature conducted at the start of the Neolnnate Study provided insight into the importance of collecting accurate delivery and labour data. We were particularly concerned that the inflammation believed to be responsible for the hypoferremic response in the neonate, may be confounded by pro- or anti-inflammatory stimuli received during labour and delivery. This included the effect of differing medication prescribed (i.e. gentamycin during labour),<sup>16</sup> and delivery methods.<sup>17</sup> As a consequence of this review, the subsequent study collected a detailed assessment of the labour process for each delivery. This study design also ensured that all recruited neonates had a cord blood sample collected using a standardised protocol, as previous studies have shown differing delays between cord-cutting can significantly affect early neonatal iron metabolism.<sup>18</sup> Importantly, it also added much to the standard of care offered to study participants.

A fundamental aspect of the study design was the allocation of all neonates to four independent groups (e.g. FTB+NBW, FTB+LBW, PTB+NBW, PTB+LBW). This has allowed for the analysis of the independent effects of both birthweight and prematurity on neonatal hypoferrremia. However, we did struggle to recruit full-term, low birthweight neonates during the study recruitment period. As birthweight is a predictor of prematurity, I believe this could be due to the research clinician, who conducted the New Ballard Score (NBS) of each neonate, not being blinded to the birthweight before NBS completion. Thus, there could be a degree of misclassification. The study research clinician was however, well trained in NBS and was deliberately unaware of specific group recruitment rates.

An important ethical issue facing the design of this study was the earliest time point in which a neonate could be bled post-delivery. Our team and hospital staff concluded that six hours would be the ideal time. This aspect of study design allowed the mother to recover physically and psychologically before providing full consent to the recruitment of the neonate and collection of the V1 sample. It also allowed clinical staff to assess if the mother or newborn were experiencing any medical complications. This led to the assessment of gestational age and anthropometrics at 6-24 hours post-delivery being a less stressful experience for the neonate, after being warmed and exposed to early breastfeeding.

Another of the most ethically complex study design aspects was the size of the blood volume collected from all recruited neonates, especially those born preterm and low birthweight. A sample volume of 3.5 ml of blood was required from each venous bleed of a full-term, normal birth weight neonate. This was calculated from the data produced by Howie *et al.*,<sup>19</sup> which suggest to the Gambia Government/MRC Joint Ethics Committee to allow a single blood draw of 2.4% total blood volume. With regards to maximum cumulative draw volume allowed, Howie *et al.*, 2011 referenced multiple institutions that allowed >10ml of whole blood from a 3 kg birth weight neonate to be taken for research purposes throughout one to two months. These volumes were deemed to provided 'minimal risk' to the neonate.

Regarding preterm neonates, venous bleed volumes used in prior studies varied according to their weight, with an average of ~2ml per venous bleed. In our study, each blood draw tube was marked with a specific volume allowed from each neonate. As another layer of protection, all neonates received a complete review of systems by a research clinician or senior study nurse before any bleed was conducted. This ensured that the subject's health was of the highest priority. Furthermore, the study was designed in order to reduce the number of bleeds for each neonate. We were aware that repetitive bleeding of healthy neonates, would be unpopular with their families and expose each neonate to repetitive episodes of stress. As a result, we allocated each full-term, normal birthweight individual into a specific study group, in order to repetitively measure iron and inflammation markers at specific ranges of time over the first week of life using a maximum of two bleeds per baby.

Bleeding neonates is notoriously difficult. At the start of the study, the study team faced the complex challenge of completing 750 neonatal peripheral bleeds. On bleeding the first study participant, we became aware that the long tubing section of the vacutainer set-up was resulting in the loss of the pressure required to complete a successful blood draw. An extended study meeting successfully overcame this hurdle, where we were able to safely adapt the blood draw equipment, while still maintaining its sterility. However, in some cases, the blood volume collected was not high enough to conduct the *ex vivo* bacterial growth assay on some serum samples. This was primarily due to the difficulty of bleeding neonates, resulting in clotting taking place before the required sample volumes could be reached. Efforts were made to miniaturise the assay; however, the reduced assay-well volume caused a breakdown in the relationship previously documented between bacterial growth and TSAT. Sadly, as a result, one of the study objectives set out in the protocol (e.g. *ex vivo* bacterial growth assay) could not be achieved.

Before the study initiation, our study team was informed that bleeding neonates in the community would be an impossible challenge, primarily due to the interference from

members of the family on the bleeding procedure. Experienced studies teams had suggested that the study design should ask the mother and their newborn to return to the maternity ward during this period. However, we were aware of experiences in other communities in The Gambia, where many neonates would have been lost to follow up if this design was adopted. With no other option available to acquire samples from discharged neonates between 24-192 hour post-delivery, we recruited an immensely experienced neonatal nurse with good local connections to conduct all community bleeds. We also worked as a study team to deliver safe and effective study protocols using the knowledge of local practices and cultural structures to drive this protocol. This resulted in 224/278 community bleeds being completed.

The Neolnnate Study suffered toward the end of the study from a lack of funds. This was primarily due to the issues relating to delays in the study initiation and the recruitment rate. The lack of funds resulted in our study being unable to examine how hypoferrremia relates to maternal iron and inflammation makers (before, during and after labour). Similarly, this also led to the inability to measure IL-6, IL-22, haptoglobin (by ELISA assay) and hemopexin in all blood samples. This analysis would have informed us as to the role that inflammation and other chaperone molecules play in the reported hypoferrremic response. The use of the Cobas 400+ biochemical analyser to detect haptoglobin concentrations in cord blood in this study, has also resulted in us not being to detect mother-neonate pairs that have experienced intrauterine infections as previously stated in Buhimschi *et al.*<sup>20</sup> If we had been able to afford haptoglobin ELISA test kits, this would not have been the case.



## 7.3 Recommendations For Future Research

### 7.3.1 Role of the Placenta in Maternal-Fetal Iron Transfer

As detailed in **Chapter 3**, sufficient iron stores are critical for a healthy pregnancy. In the placental villi, syncytiotrophoblasts uptake transferrin-bound iron from the maternal circulation via transferrin receptor 1 (TFR1) (**Chapter 3 - Figure 1**).<sup>21–23</sup> TFR1 releases iron into an acidified clathrin-coated vesicle, where it is then transported into the cytoplasm of the syncytiotrophoblast by DMT-1,<sup>23</sup> Zrt/Irt-like protein ZIP8<sup>24</sup> and ZIP14,<sup>25</sup> collectively. Ferroportin, ceruloplasmin, hephaestin, and zyklopen all assist with the transportation of iron into the fetal circulation, where it binds to fetal transferrin.<sup>22,26–28</sup> However, the regulation of maternal-placental-fetal iron transport mechanisms remains unclear. A recent publication by Sangkhae *et al.* provides additional details on the effects of iron-deficient, iron-replete and iron-overload conditions on the regulation of this process in mice, humans and human trophoblasts.<sup>29</sup>

As seen in previous publications,<sup>30–32</sup> Sangkhae *et al.* detected maternal hepcidin suppression during the pregnancy to ensure increased dietary iron intake and a raised systemic serum iron concentration.<sup>29</sup> The suppression of hepcidin at this time is believed to be regulated by several factors including erythropoiesis in the mother or fetus,<sup>32</sup> oestrogen,<sup>33</sup> and progesterone receptor membrane component-1.<sup>34</sup> Conflicting evidence now exists as to whether pregnancy-induced plasma dilution may also play a role.<sup>29,35</sup>

In instances of iron-overload, both maternal and fetal hepcidin is produced at higher quantities in order to restrict further dietary uptake and protect the fetus from oxidative damage via placental iron transfer.<sup>29,35,36</sup> However, questions remain as to whether fetal iron endowment is regulated by fetal or placental hepcidin in iron-replete or iron-deficient

conditions. Previous publications have remarked that in a state of maternal iron deficiency, the fetus could be regarded as the "perfect parasite" as human fetal iron endowment is maintained regardless of maternal iron status.<sup>21</sup> However, Sangkhae *et al.* recently recorded conflicting evidence in murine models.<sup>29</sup> Instead, iron transfer to the murine fetus under iron-deficient conditions was restricted. Decreased expression of ferroportin on the fetal side of the placenta is understood to be the cause. Interestingly, also observed was increased TFR1 expression on the maternal-facing membrane of the placental syncytiotrophoblasts. This suggests that during maternal iron deficiency, iron is held in the placenta to ensure that placental metabolic homeostasis is maintained. Sufficient iron concentration of the placenta is believed to ensure oxidative phosphorylation can continue, ensuring adequate levels of ATP are still produced.<sup>29</sup> This allows the continuation of placental protein synthesis and other critical transfer functions, protecting against placental dysfunction.

Nonetheless, these findings were only observed in the murine and *in vivo* human trophoblast models.<sup>29</sup> When similar analyses were conducted on human pregnancies, expression of TFR1 protein increased, yet ferroportin expression did not change. Sangkhae *et al.* have proposed that this may be the case due to the scarcity of severely iron deficient human mothers in their study setting.<sup>29</sup> Further research is now required to fully understand these interspecies differences, and what role placental iron regulation has on the hypoferremic response discussed here.

### **7.3.2 Potential Triggers of Early Postnatal Hypoferremia**

#### **7.3.2.1 IL-6 and the JAK/STAT3 Pathway**

The initiation of hepcidin synthesis via the actions of IL-6/JAK/STAT3 pathway has been extensively characterised.<sup>37</sup> IL-6 begins by binding to its receptor activating the formation of

a hexameric plasma-membrane signalling assembly. This is formed of IL-6, IL-6 receptor, and a gp130 subunit.<sup>38</sup> Activation of JAK2 then follows, enabling the phosphorylation of its tyrosine residues and those of subsequent downstream STAT molecules. The phosphorylated STAT molecules then dimerise, translocate to the nucleus, and target the transcription promoters of the *HAMP* gene.<sup>39</sup> Transcription of the *HAMP* gene results in the synthesis of new hepcidin proteins, which are later released into the circulation. Hepcidin then binds a single ferroportin protein on the plasma membrane of enterocytes, macrophages, and hepatocytes. This causes the endocytosis and co-degradation of both bound proteins in the lysosome.<sup>40</sup> Hence, enteric absorption of dietary iron is reduced, and sequestration of iron in macrophages causes a reduction in serum iron concentration.<sup>3</sup>

Our study has now confirmed that hepcidin-mediated neonatal hypoferremia exists in all healthy neonates (**Chapter 5**). Potential triggers for this mechanism remain unclear, as IL-6 was not analysed. However, previous research has found that inflammation (signalled by IL-6 and other cytokines) is involved in the onset and progression of human labour at term in the mother,<sup>41–48</sup> in the absence of intrauterine infection. Similarly, inflammatory markers are also detected in the cord blood of the neonate.<sup>49</sup> Previous authors have suggested that the instigating stimuli for this could involve the endocrine events of labour,<sup>47,48,50</sup> mechanical distension of the membranes and cervix (smooth muscle),<sup>41,50–53</sup> placental hypoxia and/or hypo-perfusion,<sup>50,54</sup> fetal hypoxia-acidemia,<sup>55</sup> maternal pain<sup>56</sup> or exposure to infective agents.<sup>46,48,50,57</sup> We suggest that due to the impermeability of the placenta to IL-6,<sup>49,58</sup> independent stimuli of the increased inflammatory response are taking place in the mother and newborn. This theory is supported by evidence that there are significantly lower levels of IL-6 in the newborn than the mother.<sup>59</sup> We propose that this increase in IL-6 leads to the previously reported large influx of immune cells (predominantly neutrophils) into the cervix, decidua, myometrium, chorioamniotic membranes and amniotic fluid.<sup>47,60</sup> We speculate that a similar process may occur in the neonate, with the activation of circulating neutrophils around the body at delivery. Labour is associated with neutrophilia in the early neonatal

period.<sup>61</sup> Both neonatal neutrophil activation and antigen expression correlate with the length of labour.<sup>62</sup> Decreased proportions of T cells expressing CD2, CD3 and CD4, as well as increased expression of CD16 and CD56 on natural killer cells from vaginally-delivered newborns have been reported.<sup>63</sup> This is in contrast to those born by elective caesarean section. Interestingly, another publication has documented significantly higher cord hepcidin levels in neonates born via standard vaginal delivery and second stage caesarean section than those born via elective caesarean section.<sup>64</sup> This evidence suggests that stimuli on the neonate that cause the proposed inflammation-induced hypoferremia may occur during the process of labour, rather than delivery. This is reinforced by a study by Weinschenk *et al.*,<sup>62</sup> that did not observe a difference in the increased neonatal neutrophil activation between modes of delivery after the initiation of labour. We propose that the stimuli during labour could be mechanical trauma caused by contractions in the latent and active phases of labour, or the rupture of membranes leading to the exposure of the neonate to the uterine microbiome.<sup>65</sup> Both proposed stimuli would increase in severity as the length of labour progressed. In this study, we have analysed the time between arrival to the maternity ward and time of delivery, in relation to hypoferremic response. However, as many factors are responsible for the time of arrival at the hospital before delivery, especially in LMICs with the lack of transport infrastructure and financial constraints,<sup>66</sup> it is no surprise that no significant effect was observed. Future studies measuring the length of labour accurately in relation to hypoferremia could clarify this. As to whether this effect is a deliberate action to generate the hypoferremic response is unknown. Previous authors have suggested this pro-inflammatory response may also assist in organ system transition at birth (e.g. the lungs and cytokine-induced synthesis of surfactant proteins)<sup>67</sup> and/or the activation of the immune system in the newborn.<sup>68</sup>

Since hepcidin and IL-6 are not thought to cross the placenta,<sup>58,69</sup> we have proposed that this labour-induced inflammatory response leads to the increase in hepcidin expression in the newborn and hence to the systemic sequestration of iron during and immediately after

labour/delivery has taken place. Further studies are required to confirm this, with a specific focus on the differences in iron and inflammation markers in the newborn over the first 48 hours of life between vaginally, elective caesarean and emergency caesarean section deliveries. However, this hypothetical study would be challenging due to the complex confounding clinical reasons for elective and emergency caesarean section delivery.

### **7.3.2.2 Additional Pro-Inflammatory Regulators of Hepcidin Expression**

Additional research is also required to understand the role of other pro-inflammatory regulators of hepcidin expression have on early postnatal iron homeostasis.

IL-22 is a member of the IL-10–related family of cytokines, predominately expressed by lymphocytes.<sup>70</sup> The binding of IL-22 to its receptor leads to activation of Jak1 and Tyk2, causing the phosphorylation of tyrosine molecules on STAT1, STAT3, STAT5.<sup>71</sup> The subsequent transcription of the *HAMP* gene results in the synthesis of hepcidin and the reduction of circulating serum iron levels. The distinct familial differences between IL-22 and IL-6, lead the IL-22 pathway to work independently of IL-6-mediated hepcidin expression.<sup>72</sup> IL-22 also mediates the production of other antimicrobial proteins and acute-phase reactant proteins in the liver.<sup>73</sup> These include the chaperone protein responsible for restricting bacterial uptake of human haemoglobin in the blood, haptoglobin. This finding, besides the weak activation of IL-22 from exposure to bacterial LPS,<sup>74</sup> makes additional research into this cytokine pathway necessary. This research would provide further insight as to whether contact with microorganisms during the delivery process is the exclusive trigger of neonatal hypoferremia.

Similar to IL-22, IL-1 is a pro-inflammatory molecule that can work independently of the IL-6/JAK/STAT3 signalling pathway to induce hepcidin expression.<sup>75</sup> The synthesis of hepcidin

in hepatocytes in both wild-type and IL-6 pathway knockout mice supports this observation.<sup>75</sup> Nonetheless, the action of IL-1 is suggested to more commonly be additive to the mechanism of IL-6. STAT3 and NF-κB pathway crossover is believed to be the cause.<sup>76,77</sup> This is supported by murine models showing increased hepcidin synthesis in response to LPS inoculation, compared to those that received IL-6 treatment.<sup>78</sup> IL-1 binds to its receptor, IL-1R, causing the phosphorylation of the cellular protein, IKK. Phosphorylation and degradation of pathway inhibitor, IκB then follows, resulting in the activation of NF-κB. NF-κB promotes the expression of C/EBPδ, which then binds to P-Smad and promotes the transcription of the *HAMP* gene.<sup>78</sup> IL-1Ra is a structurally similar molecule to IL-1; however, it binds to IL-1 receptor in a competitive nature and does not induce any intracellular response<sup>79</sup>. At birth, levels of IL-1Ra increase for the first days of life.<sup>80</sup> This suggests that IL-1-mediated inflammation and subsequent hepcidin synthesis may be unfavourable to early neonatal life, which is in conflict with the conclusions of our study (**Chapters 5 and 6**). An analysis of IL-1 and its relationship with the mechanisms of neonatal hypoferremia is required.

Pro-inflammatory regulators of hepcidin expression also include activin b<sup>81</sup>, leptin,<sup>82</sup> oncostatin M<sup>83,84</sup> and leukaemia inhibitory factor (LIF).<sup>84</sup> However, the strength of their independent hypoferremia-inducing qualities in the setting of bacterial infection in the human host remain unknown.

### ***7.3.2.3 Hepcidin-Independent Mechanisms of Hypoferremia***

Ferroportin (FPN1) is the only cell-surface transmembrane protein known to export ferrous iron out of mammalian cells.<sup>85</sup> Subsequently, FPN1 expression levels, play a key role in the control of cellular and systemic iron concentrations. Duodenal enterocytes, placental syncytiotrophoblasts, hepatocytes and reticuloendothelial macrophages are the predominant

cell types that express FPN1.<sup>86</sup> As seen in **Figure 1.7**, erythropoiesis, hypoxia, iron status, and inflammation are all reported to alter the expression levels of FPN1.<sup>85</sup> This is primarily controlled via the action of its master regulator, hepcidin.<sup>37</sup> Inhibition of ferroportin begins by hepcidin binding, inducing endocytosis and co-degradation of both molecules.<sup>87</sup> This results in a reduction in iron export and an increase in pooled intracellular iron.

Nonetheless, FPN1 can also be regulated via hepcidin-independent mechanisms leading to a reduction in systemic serum iron concentration.<sup>88</sup> Whether hepcidin-independent mechanisms contribute to the production of early neonatal hypoferremia (via the direct suppressive effects of inflammation on FPN1) remains undetermined.

One such potential hepcidin-independent mechanism is the activation of TLR2/6, leading to the direct downregulation of FPN1 mRNA synthesis.<sup>89,90</sup> This results in intracellular iron sequestration and systemic hypoferremia. Previous authors speculate that TLR2 and TLR6 do not activate hepcidin upregulation, offering redundancy to the mechanism of hypoferremia, enabling a faster, broader and more effective immune response.<sup>89,90</sup> These same studies have shown that bacterial lipopeptides (e.g. Mycoplasma-derived FSL1 and bacterial LPS) that target TLR2 and TLR6 promptly reduce ferroportin mRNA production for over three hours.<sup>89,90</sup> This is supported by TLR2-knockout models exhibiting no effect on ferroportin expression in response to FSL1 inoculation.<sup>89</sup> Further analyses are now required to understand if whole pathogens can trigger a similar response. This information would strengthen TLR2/6 as essential mediators of iron redistribution and may offer an alternative, more direct drug target compared to hepcidin-mediated pathways.

Tumour necrosis factor- $\alpha$  (TNF- $\alpha$ ) is a cytokine produced by many cell types, including the primary producers: macrophages, Langerhans cells, and Kupffer cells.<sup>91</sup> Following synthesis and release, TNF- $\alpha$  binds to either tumour necrosis factor receptor 1 or 2 (TNFR1 and TNFR2) resulting in the initiation of mitogen-activated protein kinase (MAP kinase), caspase,

and NF- $\kappa$ B pathways (amongst others).<sup>92</sup> In 1989, Alvarez-Hernández *et al.* proposed that TNF- $\alpha$  causes systemic hypoferrremia within the first six hours of inoculation, while total iron-binding capacity remains constant.<sup>93</sup> Laftah *et al.* uncovered a similar reduction in serum iron recorded in the first 3 hours after intraperitoneal inoculation of TNF- $\alpha$ .<sup>94</sup> These findings are supported by evidence that TNF- $\alpha$  mediated hypoferrremia is initiated within the first eight hours after caecal ligation and puncture in wild-type murine models.<sup>95</sup> No such response was detected in TNF- $\alpha$  deficient model. As no change in serum hepcidin concentration was linked to any of these responses,<sup>93–95</sup> studies suggest hepcidin-independent mechanisms regulate this method of protection, resulting in the downregulation in FPN1 transcription. TNF- $\alpha$  mediated hypoferrremia may regulate a proportion of the early neonatal hypoferrremia we have uncovered; however, additional research is needed to understand its role.

### 7.3.3 Harnessing Iron to Fight Infections

As previous research by our group has shown, limited iron concentrations reduce bacterial growth rates of sepsis-causing pathogens in human serum (**Annex 1.16**).<sup>4</sup> The NeoInnate Study discussed here has also shown that iron-sequestration is a potentially fundamental aspect of the early life innate immune response. In light of the innate and adaptive immunological differences between preterm, full-term and low birthweight newborns,<sup>96</sup> the observation that neonatal hypoferrremia is maintained in the population, regardless of gestational age or weight, suggests that it offers a significant benefit to the newborn. As a result, it is possible to consider that the limitation of iron sources to invading organisms in a multitude of other infection types could offer a partial (in combination with antibiotics) or complete (with the use of the host's immune system) treatment option. We hypothesise that limiting the growth and replication rates of invading organisms by providing a transient bacteriostatic mechanism of protection, would allow the host's neutrophils to engulf and destroy bacterial cells. We also propose that reduced replication rates of the bacterial



pathogen may allow for a reduction in the recommended dose, frequency and/or duration of current antibiotic treatments. We wonder whether combining iron-sequestration plus antibiotics may allow for toxic drug regimens and classes to be made available to individuals with a decreased maximum tolerable dose. This is particularly the case with neonates, as they possess unique physiological processes compared to adults leading to differences in drug absorption, distribution, metabolism, and elimination.<sup>97</sup>

An added benefit of this proposed treatment method would be that it is defined as combination therapy. Evidence suggests that antibiotic combination therapies already offer an increased benefit against blood infections, including those caused by carbapenem-resistant *K. pneumonia* (i.e. a common neonatal pathogen).<sup>98</sup> We speculate that similar observations may be seen by inducing transient hypoferremia, without causing long term adverse effects to iron homeostasis (**Chapter 6**). This proposed treatment strategy may allow for the use of older disused drugs to be redeployed, as observed with other drug combination therapies.<sup>99</sup> Furthermore, the use of antibiotic combination therapy has been found to reduce the spread of antimicrobial resistance.<sup>100</sup> This observation, along with most microbial pathogens requiring iron for growth and virulence, and bacterial iron acquisition mechanisms being genetically coded by pathogenic housekeeping genes, suggests that hypoferremia-antibiotic therapies may offer a reduction in the creation of new antimicrobial resistance mechanisms. Evidence of the effect that iron chelation therapy could offer in combination with an antibiotic can already be seen in the study conducted by Coraca-Huber *et al.*<sup>101</sup> This study showed that this form of treatment significantly reduced the formation of staphylococcal biofilms during infections. It is worth noting that hepcidin has also been previously described to have direct bactericidal effects on bacterial pathogens.<sup>102</sup>

The sequestration of iron and its moieties by native chaperone molecules in order to produce a protective response against invading organisms could also offer a new and novel treatment option. A study that supports this is that of Remy *et al.*,<sup>103</sup> the main finding of

which is the positive effect haptoglobin treatment has on reducing canine pneumonia and sepsis in animal models. Haptoglobin works by binding to cell-free haemoglobin in the blood and other body fluids, reducing its availability as an iron source to many bacterial pathogens.<sup>104</sup> Cell-free haemoglobin is regularly elevated during sepsis, with the level of increase correlating with a higher rate of mortality.<sup>105</sup> Remy *et al.* have shown that in a canine *S. aureus* pneumonia model with septic shock, that human haptoglobin concentrate infusions lead to the binding with canine cell-free haemoglobin. This results in its clearance and lowering of the overall level of iron within the circulation. The resulting effect is an increase in survival in the canine model. The theory behind this is an increase in the internalisation of haptoglobin-haemoglobin complexes, thus limiting access to iron for extracellular organisms. This suggests that haptoglobin therapy can enhance innate host immunity, supporting the notion it offers a novel approach to treat systemic bacterial infections. Further clinical trials in humans are required to confirm this. This use of haptoglobin is particularly relevant to neonates due to enhanced erythrocyte turnover in early postnatal life leading to the release of haemoglobin, along with the suggestion that haptoglobin synthesis is increased in response to intrauterine infection.<sup>20</sup> This supports the hypothesis it may be an effective method of innate protection.

Another treatment option in the host-pathogen battle for iron is the use of lipocalin-2. Lipocalin-2, also known as neutrophil gelatinase-associated lipocalin (NGAL), is released by several cell types (e.g. hepatocytes, pneumocytes, renal epithelial cells, and vascular smooth muscle cells) as a result of inflammation, ischemia and infection.<sup>106</sup> Recent research has shown that lipocalin-2 is produced in response to the infection of macrophages by intracellular bacterial pathogens, such as *Brucella abortus*.<sup>107</sup> Lipocalin-2 is thought to prevent iron uptake by *B. abortus* by two mechanisms: 1) by stripping iron from iron-laden siderophores and 2) intracellular levels of lipocalin-2 in macrophages inhibit further iron uptake by macrophages during infection. This results in the starvation of the pathogen of an iron source, which is critical for growth, virulence and replication. This suggests that

synthetically increasing the concentration of lipocalin-2 in the host's extracellular environment could also protect individuals against extracellular pathogens which commonly cause sepsis.<sup>108</sup>

Lactoferrin is a glycoprotein belonging to the innate immune system, found in saliva, blood, tears and human milk. Lactoferrin's central role is to bind to iron, with additional immunostimulatory, antimicrobial and anti-inflammatory roles.<sup>109</sup> Due to the structural similarities between bovine and human lactoferrin, the biological function is similar.<sup>110</sup> A Cochrane review of six studies detailing its use in preterm neonates has shown that oral supplementation of lactoferrin with or without probiotics reduced the risk of late-onset necrotizing enterocolitis, bacterial and fungal sepsis.<sup>111</sup> Equally, the use of recombinant lactoferrin (i.e. Talactoferrin) offers some promise as it does not require pasteurisation, increasing its biological activity.<sup>112</sup> Interestingly, concentrations of lactoferrin are high in colostrum,<sup>113</sup> again suggesting that iron-restriction is an essential aspect of neonatal innate immune defence in the gut.<sup>114</sup> How this might be linked to circulating levels of neonatal iron remains unclear.

Lastly, research is also progressing toward harnessing the use of the bacterium's iron uptake machinery in order to combat the infections it causes. This comes in the form of the manipulation of the action of bacterial siderophores and the bacterial influx transport channels. Small-molecule siderophores are produced and released by nearly all bacterial species.<sup>115</sup> As seen in **Figure 1.6** and **Table 1.2**, the redundancy of these iron-uptake systems allows bacteria to bind and consume different forms of iron. This function of multiple ligand targets by a plethora of siderophore types is predominately due to their differing chemical structures of the iron-chelating chemical groups (i.e. hydroxamate,  $\alpha$ -hydroxyl-carboxylate and catechol).<sup>115</sup> The ability to synthesise these molecules is often transferred between bacterial isolates by mobile genetic elements.<sup>116</sup> Recently, developments have been made to exploit these iron-siderophore uptake systems in common bacterial pathogens

by using siderophores conjugated to antibiotics. This new form of antibiotics has been focused on gram-negative bacteria due to the threat they pose with regards to antimicrobial resistance and their vastly characterised bacterial iron uptake mechanisms.<sup>117</sup> One drug discovery candidate that is showing early promise is cefiderocol. The new expedited development program of cefiderocol has led to a recent international, double-blind, randomised phase III clinical trial of the efficacy and safety of cefiderocol in patients with nosocomial pneumonia.<sup>118</sup> These results highlight that cefiderocol met the non-inferiority comparison to high dose meropenem in all-cause mortality at 14 days after initiation of treatment. Cefiderocol, a siderophore-cephalosporin drug, which uses the bacterium's active transport machinery in a "Trojan horse" style to gain entry past the cell wall. As previously stated, cefiderocol is now of particular use against carbapenem-resistant gram-negative pathogens, including those expressing New Delhi metallo- $\beta$ -lactamase-mediated carbapenem resistance, as seen in cases of neonatal sepsis (**Section 1.9.7**). Additionally, *in vitro* activity against isolates from the global surveillance studies, SIDERO-WT and SIDERO-CR have shown that cefiderocol is highly effective against current strains of *P. aeruginosa*, *A. baumannii*, *B. cepacia*, *B. pseudomallei* and *S. maltophilia*.<sup>119,120</sup> Several of these have recently been noted in cases of invasive bacterial infection in neonates in sub-Saharan Africa.<sup>121</sup>

### 7.3.4 Hecpidin Agonists as Hypoferremic Therapies

Here, we speculate that the use of hepcidin agonists or inducers may be the hypoferremic therapy that is required to produce this iron-restricted environment in the host. There are several therapeutic inducers of hepcidin expression; these include BMP6, TMPRSS6-silencing oligonucleotides and a number of small molecules.<sup>122</sup> One of the inducers that has progressed in its development is IONIS-TMPRSS6-LRX (Ionis Pharmaceuticals, Inc.).<sup>123</sup> This molecule is in phase 2 clinical trials and induces hepcidin expression by silencing

Tmprss6 mRNA by using lipid nanoparticles linked to small interfering RNAs (siRNAs) (ClinicalTrials.gov Identifier: NCT04059406).

A hepcidin agonist of note is minihepcidin PR73, which works by inhibiting the function of the transmembrane protein, ferroportin. PR73 has been shown to have possible protective effects in animal models against systemic bacterial infection. Researchers have uncovered that animal models deficient in hepcidin, when given PR73, are protected against siderophilic bacteria such as *Vibrio vulnificus*<sup>1</sup> and *Yersinia enterocolitica*.<sup>124</sup> Additionally, PR73 is observed to be protective against non-siderophilic bacteria such as *Klebsiella pneumoniae* or *Escherichia coli*.<sup>125,126</sup> This is due to the PR73 mechanism of action leading to a systemic hypoferremic state in the model, repressing bacterial growth, and subsequently reducing the likelihood of mortality. These findings are especially remarkable as a similar deficiency of hepcidin, seen in hereditary hemochromatosis, leads to increased mortality when infected with siderophilic bacteria.<sup>127,128</sup> This suggests that these observations may be relatable to humans. Currently, all of the research relating to the use of PR73 is in the preclinical phase.<sup>122</sup>

Two hepcidin agonists that have reached clinical trials are LJPC-401 (La Jolla Pharmaceutical Company) and PTG-300 (Protagonist Therapeutics Inc.). PTG-300 is now in phase II and has previously been shown to reduce serum iron concentration in cynomolgus monkeys.<sup>129</sup> During phase I trials, the drug was shown to be tolerated by study participants and reduced their serum iron concentration in a dose-dependent manner. Hypoferremic responses were maintained for almost 144 hours.<sup>130</sup> The LJPC-401 phase 1 trial reports similar findings with reduced serum iron concentration of the blood in a dose-dependent manner. Researchers found that the maximal point of hypoferremia occurred at eight-hours post-injection.<sup>131</sup>

Though the clinical trials referenced suggest that induced hepcidin levels produce a fast-acting and short-lived response, it is still unclear what effect hormonal augmentation would have in the sick neonate in the immediate period after treatment, or the long-term effects (i.e. erythropoiesis, microbiome and iron deficiency). Further research is required in humans to assess how we might produce a balanced therapy, focused on 1) maintaining iron-sequestration for a required time allowing for iron starvation in invading organisms, and 2) to not produce significant adverse side effects to human iron homeostasis. Realistically it will be many years before trials in neonates would be seriously contemplated.

### 7.3.5 Hepcidin-Resistance

Assessing our data on iron and inflammation markers in full-term, normal birthweight neonates over the first week of life (**Chapter 6**) leads us to question the validity of the proposed methods of treatment for severe bacterial infections in newborns seen in **Sections 7.3.3 and 7.3.4**. The hepcidin-induced hypoferremia observed in the Neonate Study was a robust and fast-acting response to early postpartum life. Nevertheless, it was not maintained for longer than 72 hours of life in most individuals. This is even the case despite hepcidin levels remaining higher than adult levels in the majority of the full term, normal birthweight study participants over the first week of life. From our data, we suggest that hepcidin-resistance may potentially halt the continuation of this protective mechanism after 48 hours post-delivery. This is observed in **Chapter 6 (Figures 3 and 4)** where hepcidin levels are maintained at high levels at 48 hours onwards, with serum iron and TSAT levels increasing over the same period. We hypothesise that this is due to the effect of high dose excess hepcidin, causing occlusion rather than degradation of ferroportin on the membrane of neonatal macrophages. Previous research has shown that excess hepcidin can preferentially cause occlusion, leading to the intermittent binding and releasing of hepcidin to the central cavity of individual ferroportin molecules.<sup>132</sup> We propose that this may allow for

the steady and slow increase of serum iron concentration over the following days. This finding is noteworthy, as dietary iron absorption during this period is very low.<sup>133</sup> Therefore, any increase in serum iron is unlikely to be caused due to enterocyte iron efflux.

A previous study in suckling 15-day old mouse models has found a potentially related discovery in the context of early life hepcidin resistance.<sup>134</sup> Frazer *et al.* aimed to elicit if, as previous studies had suggested, iron absorption in suckling mammals was resilient to stimuli that ordinarily decreased absorption in adults. Interestingly, this study uncovered that enterocyte ferroportin was hyporesponsive to the effect of circulating serum hepcidin during the suckling period. Immunofluorescence assays suggested that this finding was not due to changes in ferroportin localisation, making it inaccessible to the actions of hepcidin, as previously reported by Theurl *et al.* concerning hepcidin resistance in the retinal pigment epithelium.<sup>135</sup> Similarly, this could be the possible cause of hepcidin-resistance in our study, due to impermeable physical barriers between molecules of ferroportin and hepcidin in a subset of cell types responsible for iron release at the latter part of the first week of life. On balance, this is unlikely to be the case as we believe circulating and splenic macrophages are the target cells for iron sequestration during neonatal hypoferremia.

Furthermore, Frazer *et al.* found that decreases in serum iron levels occurred after hepcidin stimulation, suggesting that hepcidin activity was present in some cell types.<sup>134</sup> However, this was not the case concerning enterocytes. After further laboratory analysis, ferroportin in these murine enterocytes was shown to be smaller than standard adult ferroportin molecules. The authors suggested that this may be due to early life murine ferroportin being spliced or glycosylated differently. This evidence suggests that in murine models, age-specific modification of ferroportin is found in the transmembrane of enterocytes. Further research is required to assess if that is the case with respect to human neonatal macrophage-expressed ferroportin molecules.

Similar to our reports, Frazer *et al.* note that hepcidin-resistance could be the result of high hepcidin excess favouring the occlusion of ferroportin rather than its degradation. As seen in Aschemeyer *et al.* hepcidin is thought to bind to the central metal-binding cavity, leading to the hinderance of the conformational changes required for iron efflux.<sup>132</sup> Additionally, we submit that with the hepcidin levels in excess, intermittent binding and releasing of hepcidin to the central cavity of individual ferroportin molecules may allow for slow rates of iron efflux into the circulation (**Chapter 6**). This leads us to consider that there could be scope for research into the optimum dose of minihepcidin required to swing the balance back into the favour of degradation rather than occlusion, should our hypothesis be found to be correct. Equally, it would also be interesting to access the endocytic machinery of neonatal macrophages to uncover if deficiencies in degradation mechanisms may be the cause of these proposed reasons for hepcidin-resistance.<sup>136,137</sup>

Why hepcidin-resistance occurs in young mice and humans remains unclear. However, it could be that both the hyporesponsiveness of murine enterocytes in 15-day mice pups and the hepcidin-resistance uncovered in our study in human neonates is a semi-protective mechanism against the iron deficiency in early life, which is particularly common.<sup>138</sup> It may be possible that human neonates are trying to strike a balance between inducing protective hypoferremia at the point it is most required to protect against infection (i.e. 24 hours of life) and ensuring serum iron is still available to the tissues responsible for erythropoiesis and growth.

Further research is required to determine whether neonatal ferroportin is differentially spliced or glycosylated and if so, what cell types this occurs in. It is also crucial that we establish whether, like adult ferroportin, the expression of the hypothetical variant form of ferroportin suggested here, is less or more affected by inflammation and by what mechanism.



Until these questions are answered, it would not be worthwhile supporting the use of hepcidin agonist or inducers to combat neonatal systemic infections.

## **7.4 Conclusions**

The research of this thesis has contributed three key components to our present understanding of neonatal iron homeostasis. Firstly, it has evaluated our current knowledge of iron and inflammation markers (hepcidin, serum iron and TSAT) in the neonatal period in respect to gestational age. This was only possible because of the newly developed methods in the standardisation of hepcidin values across multiple retrospective studies in order to produce comparisons and averages. Secondly, it has unequivocally refuted our previous hypothesis that immature or growth-restricted neonates might have a lesser ability to trigger a hypoferremic defence and that this might explain their greater susceptibility to septicaemias. Consequently, we have shown that the premature and low birthweight neonates all exhibited a profound hypoferremia during the first 24 hours of life, with no detectable differences from the full-term, normal birthweight controls. The remarkable similarity in the hypoferremic response across all the study groups further accentuates the efficiency of the process and supports the likelihood that it occurs by the process of evolution. Thirdly, our results suggest that after a fast-acting, short-lived early postnatal hypoferremic response at birth, there is a period of hepcidin-resistance, resulting in slow and consistent iron efflux into the circulation over the next week of life. Further research is required to understand why neonatal iron homeostasis is unresponsive to high serum hepcidin concentrations after the hypoferremic period. While contemplating the use of iron-sequestration as a method of combatting antimicrobial resistant infections seems untimely, in light of our results and the threat we all face, the importance of scrutinising hypoferremia as a mechanism of protection has never been more evident.

## 7.5 References

- 1 Arezes J, Jung G, Gabayan V, *et al.* Hepcidin-induced hypoferremia is a critical host defense mechanism against the siderophilic bacterium *Vibrio vulnificus*. *Cell Host Microbe* 2015; **17**: 47–57.
- 2 Prentice S, Jallow AT, Sinjanka E, *et al.* Hepcidin mediates hypoferremia and reduces the growth potential of bacteria in the immediate post-natal period in human neonates. *Sci Rep* 2019; **9**: 16596
- 3 Ganz T, Nemeth E. Iron Sequestration and Anemia of Inflammation. *Semin Hematol* 2009; **46**: 387–93.
- 4 Cross JH, Bradbury RS, Fulford AJ, *et al.* Oral iron acutely elevates bacterial growth in human serum. *Sci Rep* 2015; **5**: 16670.
- 5 Balogh A, Szabo M, Kelen D, *et al.* Prohepcidin levels during human perinatal adaptation. *Pediatr Hematol Oncol* 2007; **24**: 361–8.
- 6 Kitajima J, Ohga S, Kinjo T, *et al.* Serum prohepcidin concentrations at birth and 1 month after birth in premature infants. *Pediatr Blood Cancer* 2011; **56**: 267–72.
- 7 Yamada RRT, Leone CCR. Hematological and iron content evolution in exclusively breastfed late-preterm newborns. *Clinics* 2014; **69**: 792–8.
- 8 Szabo M, Vasarhelyi B, Balla G, *et al.* Acute postnatal increase of extracellular antioxidant defence of neonates: the role of iron metabolism. *Acta Paediatr* 2001; **90**: 1167–70.
- 9 Wang L, Xu J, Tian Y, Liu Y, Wu H. [Expression and significance of Toll-like receptors in cord blood mononuclear cells.]. *Chinese J Pediatr* 2007; **45**: 365–8.
- 10 Cross JH, Jarjou O, Mohammed NI, Prentice AM, Cerami C. Neonatal iron distribution and infection susceptibility in full term, preterm and low birthweight babies in urban Gambia: study protocol for an observational study. *Gates Open Res* 2019; **3**: 1469.
- 11 van der Vorm LN, Hendriks JCM, Laarakkers CM, *et al.* Toward Worldwide Hepcidin

- Assay Harmonization: Identification of a Commutable Secondary Reference Material. *Clin Chem* 2016; **62**: 993–1001.
- 12 Unger H, Thriemer K, Ley B, *et al.* The assessment of gestational age: a comparison of different methods from a malaria pregnancy cohort in sub-Saharan Africa. *BMC Pregnancy Childbirth* 2019; **19**: 12.
  - 13 Mbofung CMF, Atinmo T, Omolulu A. Mineral content of colostrum and mature milk of lactating Nigerian women as influenced by stage of lactation. *Nutr Rep Int* 1984; **30**: 1137–46.
  - 14 Lee S, Guillet R, Cooper EM, *et al.* Prevalence of anemia and associations between neonatal iron status, hepcidin, and maternal iron status among neonates born to pregnant adolescents. *Pediatr Res* 2016; **79**: 42–8.
  - 15 Basu S, Kumar N, Srivastava R, Kumar A. Maternal and Cord Blood Hepcidin Concentrations in Severe Iron Deficiency Anemia. *Pediatr Neonatol* 2016; **57**: 413–9.
  - 16 Umeki S. Anti-inflammatory action of gentamycin through inhibitory effect on neutrophil NADPH oxidase activity. *Comp Biochem Physiol.* 1995; **110**: 817–21.
  - 17 Locatelli A, Pintucci A. Vacuum-assisted vaginal delivery. Operative Obstetrics, Fouth Edition. CRC Press, 2017: 317–32.
  - 18 WHO | Guideline: Delayed Umbilical Cord Clamping for Improved Maternal and Infant Health and Nutrition Outcomes. Geneva: World Health Organization; 2014. Available from: <https://www.ncbi.nlm.nih.gov/books/NBK310511/>.
  - 19 Howie SRC. Les volumes des échantillons sanguins dans la recherche en matière de santé Infantile: Examen des limites de sécurité. *Bull. World Health Organ.* 2011; **89**: 46–53.
  - 20 Buhimschi CS, Bhandari V, Dulay AT, *et al.* Proteomics mapping of cord blood identifies haptoglobin ‘switch-on’ pattern as biomarker of early-onset neonatal sepsis in preterm newborns. *PLoS One* 2011; **6**: e26111.
  - 21 Cao C, Fleming MD. The placenta: the forgotten essential organ of iron transport. *Nutr Rev* 2016; **74**: 421–31.

- 22 Bastin J, Drakesmith H, Rees M, Sargent I, Townsend A. Localisation of proteins of iron metabolism in the human placenta and liver. *Br J Haematol* 2006; **134**: 532–43.
- 23 Georgieff MK, Wobken JK, Welle J, Burdo JR, Connor JR. Identification and Localization of Divalent Metal Transporter-1 (DMT-1) in Term Human Placenta. *Placenta* 2000; **21**: 799–804.
- 24 Gálvez-Peralta M, He L, Jorge-Nebert LF, *et al.* Zip8 zinc transporter: Indispensable role for both multiple-organ organogenesis and hematopoiesis in utero. *PLoS One* 2012; **7**: e36055.
- 25 Hojyo S, Fukada T, Shimoda S, *et al.* The zinc transporter SLC39A14/ZIP14 controls G-protein coupled receptor-mediated signaling required for systemic growth. *PLoS One* 2011; **6**: e18059.
- 26 Chen H, Attieh ZK, Syed BA, *et al.* Identification of zyklopen, a new member of the vertebrate multicopper ferroxidase family, and characterization in rodents and human cells. *J Nutr* 2010; **140**: 1728–35.
- 27 Guller S, Buhimschi CS, Ma YY, *et al.* Placental expression of ceruloplasmin in pregnancies complicated by severe preeclampsia. *Lab Invest* 2008; **88**: 1057–67.
- 28 Li Y-Q, Bai B, Cao X-X, Yan H, Zhuang G-H. Ferroportin 1 and hephaestin expression in BeWo cell line with different iron treatment. *Cell Biochem Funct* 2012; **30**: 249–55.
- 29 Sangkhae V, Fisher AL, Wong S, *et al.* Effects of maternal iron status on placental and fetal iron homeostasis. *J Clin Invest* 2020; **130**: 625-640.
- 30 Rehu M, Punnonen K, Ostland V, *et al.* Maternal serum hepcidin is low at term and independent of cord blood iron status. *Eur J Haematol* 2010; **85**: 345–52.
- 31 Kulik-Rechberger B, Kościeszka A, Szponar E, Domsud J. Hepcidin and iron status in pregnant women and full-term newborns in first days of life. *Ginekol Pol.* 2016; **87**:288-92.
- 32 Koenig M, Tussing-Humphreys L, Day J, Cadwell B, Nemeth E. Hepcidin and Iron Homeostasis during Pregnancy. *Nutrients* 2014; **6**: 3062–83.
- 33 Hou Y, Zhang S, Wang L, *et al.* Estrogen regulates iron homeostasis through

- governing hepatic hepcidin expression via an estrogen response element. *Gene* 2012; **511**: 398–403.
- 34 Li X, Rhee DK, Malhotra R, *et al.* Progesterone receptor membrane component-1 regulates hepcidin biosynthesis. *J Clin Invest* 2015; **126**: 389–401.
  - 35 Fisher AL, Nemeth E. Iron homeostasis during pregnancy. *Am J Clin Nutr* 2017; **106**: 1567S-1574S.
  - 36 Nicolas G, Bennoun M, Porteu A, *et al.* Severe iron deficiency anemia in transgenic mice expressing liver hepcidin. *Proc Natl Acad Sci* 2002; **99**: 4596–601.
  - 37 Rishi G, Wallace DF, Subramaniam VN. Hepcidin: Regulation of the master iron regulator. *Biosci. Rep.* 2015; **35**: e00192.
  - 38 Lacroix M, Rousseau F, Guilhot F, *et al.* Novel insights into interleukin 6 (IL-6) Cis- And transsignaling pathways by differentially manipulating the assembly of the IL-6 signaling complex. *J Biol Chem* 2015; **290**: 26943-53.
  - 39 Wrighting DM, Andrews NC. Interleukin-6 induces hepcidin expression through STAT3. *Blood* 2006; **108**: 3204–9.
  - 40 Preza GC, Pinon R, Ganz T, Nemeth E. Cellular Catabolism of the Iron-Regulatory Peptide Hormone Hepcidin. *PLoS One* 2013; **8**: e58934.
  - 41 Santhanam U, Avila C, Romero R, *et al.* Cytokines in normal and abnormal parturition: elevated amniotic fluid interleukin-6 levels in women with premature rupture of membranes associated with intrauterine infection. *Cytokine* 1991; **3**: 155–63.
  - 42 Opsjøn S-L, Wathen NC, Tingulstad S, *et al.* Tumor necrosis factor, interleukin-1, and interleukin-6 in normal human pregnancy. *Am J Obstet Gynecol* 1993; **169**: 397–404.
  - 43 Liechty KW, Koenig JM, Mitchell MD, Romero R, Christensen RD. Production of interleukin-6 by fetal and maternal cells in vivo during intraamniotic infection and in vitro after stimulation with interleukin. *Pediatr Res* 1991; **29**: 1–4.
  - 44 Hillier SL, Witkin SS, Krohn MA, Watts DH, Kiviat NB, Eschenbach DA. The relationship of amniotic fluid cytokines and preterm delivery, amniotic fluid infection,

- histologic chorioamnionitis, and chorioamnion infection. *Obstet Gynecol* 1993; **81**: 941–8.
- 45 Austgulen R, Lien E, Liabakk NB, Jacobsen G, Arntzen KJ. Increased levels of cytokines and cytokine activity modifiers in normal pregnancy. *Eur J Obstet Gynecol Reprod Biol* 1994; **57**: 149–55.
  - 46 Cox SM, Casey ML, MacDonald PC. Accumulation of interleukin-1 $\beta$  and interleukin-6 in amniotic fluid: A sequela of labour at term and preterm. *Hum. Reprod. Update*. 1997; **3**: 517–27.
  - 47 Christiaens I, Zaragoza DB, Guilbert L, Robertson SA, Mitchell BF, Olson DM. Inflammatory processes in preterm and term parturition. *J. Reprod. Immunol.* 2008; **79**: 50–7.
  - 48 Keelan JA, Blumenstein M, Helliwell RJA, Sato TA, Marvin KW, Mitchell MD. Cytokines, prostaglandins and parturition - A review. *Placenta* 2003; **24**: S33-46.
  - 49 Duncombe G, Veldhuizen RAW, Gratton RJ, Han VKM, Richardson BS. IL-6 and TNF $\alpha$  across the umbilical circulation in term pregnancies: Relationship with labour events. *Early Hum Dev* 2010; **86**: 113–7.
  - 50 Bowen JM, Chamley L, Mitchell MD, Keelan JA. Cytokines of the placenta and extra-placental membranes: Biosynthesis, secretion and roles in establishment of pregnancy in women. *Placenta*. 2002; **23**: 239–56.
  - 51 El Maradny E, Kanayama N, Halim A, Maehara K, Terao T. Stretching of fetal membranes increases the concentration of interleukin- 8 and collagenase activity. *Am J Obstet Gynecol* 1996; **174**: 843–9.
  - 52 Maehara K, Kanayama N, Maradny EE, Uezato T, Fujita M, Terao T. Mechanical stretching induces interleukin-8 gene expression in fetal membranes: a possible role for the initiation of human parturition. *Eur J Obstet Gynecol Reprod Biol* 1996; **70**: 191–6.
  - 53 Sivarajasingam SP, Imami N, Johnson MR. Myometrial cytokines and their role in the onset of labour. *J. Endocrinol.* 2016; **231**: R101–19.

- 54 Pierce BT, Pierce LM, Wagner RK, *et al.* Hypoperfusion causes increased production of interleukin 6 and tumor necrosis factor  $\alpha$  in the isolated, dually perfused placental cotyledon. *Am J Obstet Gynecol* 2000; **183**: 863–7.
- 55 Chiesa C, Pellegrini G, Panero A, *et al.* Umbilical cord interleukin-6 levels are elevated in term neonates with perinatal asphyxia. *Eur J Clin Invest* 2003; **33**: 352–8.
- 56 Saito S, Kasahara T, Kato Y, Ishihara Y, Ichijo M. Elevation of amniotic fluid interleukin 6 (IL-6), IL-8 and granulocyte colony stimulating factor (G-CSF) in term and preterm parturition. *Cytokine* 1993; **5**: 81–8.
- 57 Gotsch F, Romero R, Kusanovic JP, *et al.* The fetal inflammatory response syndrome. *Clin. Obstet. Gynecol.* 2007; **50**: 652–83.
- 58 Aaltonen R, Heikkinen T, Hakala K, Laine K, Alanen A. Transfer of Proinflammatory Cytokines Across Term Placenta. *Obstet Gynecol* 2005; **106**: 802–7.
- 59 Opsjon SL, Austgulen R, Waage A. Interleukin-1, interleukin-6 and tumor necrosis factor at delivery in preeclamptic disorders. *Acta Obstet Gynecol Scand* 1995; **74**: 19–26.
- 60 Osman I, Young A, Ledingham MA, *et al.* Leukocyte density and pro-inflammatory cytokine expression in human fetal membranes, decidua, cervix and myometrium before and during labour at term. *Mol. Hum. Reprod.* 2003; **9**: 41–5.
- 61 Barak Yigal, Blachar Yoram, Levin Stanley. Neonatal Neutrophilia: Possible Role of a Humoral Granulopoietic Factor. *Ped Res* 1980; **14**: 1026-1028.
- 62 Weinschenk NP, Farina A, Bianchi DW. Neonatal neutrophil activation is a function of labor length in preterm infants. *Pediatr Res* 1998; **44**: 942–5.
- 63 Samelson R, Larkey DM, Amankwah Ks, McConnachie P. Effect of Labor on Lymphocyte Subsets in Full-Term Neonates. *Am J Reprod Immunol* 1992; **28**: 71–3.
- 64 Lorenz L, Herbst J, Engel C, *et al.* Gestational Age-Specific Reference Ranges of Hepcidin in Cord Blood. *Neonatology* 2014; **106**: 133–9.
- 65 Baker JM, Chase DM, Herbst-Kralovetz MM. Uterine microbiota: Residents, tourists, or invaders? *Front. Immunol.* 2018; **9**: 208.

- 66 Treacy L, Bolkan HA, Sagbakken M. Distance, accessibility and costs. Decision-making during childbirth in rural Sierra Leone: A qualitative study. *PLoS One* 2018; **13**: e0188280.
- 67 Moss TJM, Newnham JP, Willett KE, Kramer BW, Jobe AH, Ikegami M. Early gestational intra-amniotic endotoxin: Lung function, surfactant, and morphometry. *Am J Respir Crit Care Med* 2002; **165**: 805–11.
- 68 Protonotariou E, Chrelias C, Kassanos D, Kapsambeli H, Trakakis E, Sarandakou A. Immune response parameters during labor and early neonatal life. *In Vivo* 2010; **24**: 117–24.
- 69 Hoppe M, Hulthén L, Samuelson G. Is cord blood hepcidin influenced by the low-grade acute-phase response occurring during delivery? A small-scale longitudinal study. *J. Matern. Neonatal Med.* 2018; **32**: 2166-2172.
- 70 Dudakov JA, Hanash AM, van den Brink MRM. Interleukin-22: Immunobiology and Pathology. *Annu Rev Immunol* 2015; **33**: 747-785.
- 71 Lejeune D, Dumoutier L, Constantinescu S, Kruijer W, Schuringa JJ, Renauld JC. Interleukin-22 (IL-22) activates the JAK/STAT, ERK, JNK, and p38 MAP kinase pathways in a rat hepatoma cell line: Pathways that are shared with and distinct from IL-10. *J Biol Chem* 2002; **277**: 33676–82.
- 72 Smith CL, Arvedson TL, Cooke KS, *et al.* IL-22 regulates iron availability in vivo through the induction of hepcidin. *J Immunol* 2013; **191**: 1845–55.
- 73 Veas F, Dubois G. IL-22 Induces an Acute-Phase Response Associated to a Cohort of Acute Phase Proteins and Antimicrobial Peptides as Players of Homeostasis. In: *Acute Phase Proteins - Regulation and Functions of Acute Phase Proteins. Intech Open.* 2011: 20386.
- 74 Wallace DF, Subramaniam VN. Analysis of IL-22 contribution to hepcidin induction and hypoferremia during the response to LPS in vivo. *Int Immunol* 2015; **27**: 281-7.
- 75 Lee P, Peng H, Gelbart T, Wang L, Beutler E. Regulation of hepcidin transcription by interleukin-1 and interleukin-6. *Proc Natl Acad Sci U S A* 2005; **102**: 1906-10.



- 76 Bode JG, Albrecht U, Häussinger D, Heinrich PC, Schaper F. Hepatic acute phase proteins - Regulation by IL-6- and IL-1-type cytokines involving STAT3 and its crosstalk with NF- $\kappa$ B-dependent signaling. *Eur. J. Cell Biol.* 2012; 91: 496-505.
- 77 Goldstein I, Paakinaho V, Baek S, Sung MH, Hager GL. Synergistic gene expression during the acute phase response is characterized by transcription factor assisted loading. *Nat Commun* 2017; **8**: 1849.
- 78 Litwack G. Iron Metabolism : Hepcidin, 1st edn. 2019.
- 79 Arend WP, Malyak M, Guthridge CJ, Gabay C. Interleukin-1 Receptor Antagonist : Role in Biology. *Annu Rev Immunol* 1998; **16**: 27–55.
- 80 Lusyati S, Hulzebos C V., Zandvoort J, Sauer PJJ. Levels of 25 cytokines in the first seven days of life in newborn infants. *BMC Res Notes* 2013; **6**: 547.
- 81 Besson-Fournier C, Latour C, Kautz L, *et al.* Induction of activin B by inflammatory stimuli up-regulates expression of the iron-regulatory peptide hepcidin through Smad1/5/8 signaling. *Blood* 2012; **120**: 431-9.
- 82 Chung B, Matak P, McKie AT, Sharp P. Leptin Increases the Expression of the Iron Regulatory Hormone Hepcidin in HuH7 Human Hepatoma Cells. *J Nutr* 2007; **137**: 2366-70.
- 83 Chung B, Verdier F, Matak P, Deschemin J, Mayeux P, Vaulont S. Oncostatin M is a potent inducer of hepcidin, the iron regulatory hormone. *FASEB J* 2010; **24**: 2093–103.
- 84 Kanda J, Uchiyama T, Tomosugi N, Higuchi M, Uchiyama T, Kawabata H. Oncostatin M and leukemia inhibitory factor increase hepcidin expression in hepatoma cell lines. *Int J Hematol* 2009; **90**: 545–52.
- 85 Ward DM, Kaplan J. Ferroportin-mediated iron transport: Expression and regulation. *Biochim. Biophys. Acta - Mol. Cell Res.* 2012; **1823**: 1426–33.
- 86 Drakesmith H, Nemeth E, Ganz T. Ironing out Ferroportin. *Cell Metab.* 2015; **22**: 777–87.
- 87 Nemeth E, Tuttle MS, Powelson J, *et al.* Hepcidin regulates cellular iron efflux by

- binding to ferroportin and inducing its internalization. *Science* 2004; **306**: 2090–3.
- 88 Enculescu M, Metzendorf C, Sparla R, *et al.* Modelling Systemic Iron Regulation during Dietary Iron Overload and Acute Inflammation: Role of Hepcidin-Independent Mechanisms. *PLoS Comput Biol* 2017; **13**: e1005322.
  - 89 Guida C, Altamura S, Klein FA, *et al.* A novel inflammatory pathway mediating rapid hepcidin-independent hypoferremia. *Blood* 2015; **125**: 2265–75.
  - 90 Abreu R, Quinn F, Giri PK. Role of the hepcidin-ferroportin axis in pathogen-mediated intracellular iron sequestration in human phagocytic cells. *Blood Adv* 2018; **2**: 1089-1100.
  - 91 Parameswaran N, Patial S. Tumor necrosis factor- $\alpha$  signaling in macrophages. *Crit. Rev. Eukaryot. Gene Expr.* 2010; **20**: 87-103.
  - 92 Sabio G, Davis RJ. TNF and MAP kinase signalling pathways. *Semin. Immunol.* 2014; **26**: 237-245.
  - 93 Alvarez-Hernandez X, Liceaga J, McKay IC, Brock JH. Induction of hypoferremia and modulation of macrophage iron metabolism by tumor necrosis factor. *Lab Investig* 1989.
  - 94 Laftah AH, Sharma N, Brookes MJ, *et al.* Tumour necrosis factor  $\alpha$  causes hypoferraemia and reduced intestinal iron absorption in mice. *Biochem J* 2006; **387**: 61-7.
  - 95 Schubert TEO, Bosserhoff AK, Peyssonaux C, *et al.* Hypoferraemia during the early inflammatory response is dependent on tumour necrosis factor activity in a murine model of protracted peritonitis. *Mol Med Rep* 2012; **6**: 838-42.
  - 96 Melville JM, Moss TJM. The immune consequences of preterm birth. *Front. Neurosci.* 2013; **7**: 79.
  - 97 Ku LC, Brian Smith P. Dosing in neonates: Special considerations in physiology and trial design. *Pediatr. Res.* 2015; **77**: 2–9.
  - 98 Morrill HJ, Pogue JM, Kaye KS, LaPlante KL. Treatment options for carbapenem-resistant Enterobacteriaceae infections. *Open Forum Infect Dis* 2015; **2**: ofv050.

- 99 Coates A. The future of antibiotics lies in combination treatments. *Futur Drug Discov* 2019; **1**: FDD5.
- 100 Singh N, Yeh PJ. Suppressive drug combinations and their potential to combat antibiotic resistance. *J Antibiot* 2017; **70**: 1033–42.
- 101 Coraça-Huber DC, Dichtl S, Steixner S, Nogler M, Weiss G. Iron chelation destabilizes bacterial biofilms and potentiates the antimicrobial activity of antibiotics against coagulase-negative Staphylococci. *Pathog Dis* 2018; **76**: 1093.
- 102 Jiang XF, Liu ZF, Lin AF, Xiang LX, Shao JZ. Coordination of Bactericidal and Iron Regulatory Functions of Heparin in Innate Antimicrobial Immunity in a Zebrafish Model. *Sci Rep* 2017; **7**: 4265.
- 103 Remy KE, Cortés-Puch I, Solomon SB, *et al.* Haptoglobin improves shock, lung injury, and survival in canine pneumonia. *JCI insight* 2018; **3**: e123013.
- 104 Parrow NL, Fleming RE, Minnick MF. Sequestration and Scavenging of Iron in Infection. *Infect Immun* 2013; **81**: 3503–14.
- 105 Adamzik M, Hamburger T, Petrat F, Peters J, de Groot H, Hartmann M. Free hemoglobin concentration in severe sepsis: methods of measurement and prediction of outcome. *Crit Care* 2012; **16**: R125.
- 106 Chakraborty S, Kaur S, Guha S, Batra SK. The multifaceted roles of neutrophil gelatinase associated lipocalin (NGAL) in inflammation and cancer. *Biochim. Biophys. Acta - Rev. Cancer*. 2012; **1826**: 129–69.
- 107 Hop HT, Arayan LT, Huy TXN, *et al.* Lipocalin 2 (Lcn2) interferes with iron uptake by *Brucella abortus* and dampens immunoregulation during infection of RAW 264.7 macrophages. *Cell Microbiol* 2018; **20**: 12813.
- 108 Lu F, Inoue K, Kato J, Minamishima S, Morisaki H. Functions and regulation of lipocalin-2 in gut-origin sepsis: a narrative review. *Crit Care* 2019; **23**: 269.
- 109 Actor JK, Hwang S-A, Kruzel ML. Lactoferrin as a natural immune modulator. *Curr Pharm Des* 2009; **15**: 1956–73.
- 110 Baker EN, Anderson BF, Baker HM, *et al.* Structure, function and flexibility of human

- lactoferrin. *Int J Biol Macromol* 1991; **13**: 122–9.
- 111 Pammi M, Suresh G. Enteral lactoferrin supplementation for prevention of sepsis and necrotizing enterocolitis in preterm infants. *Cochrane Database Syst. Rev* 2017; **6**: CD007137.
  - 112 Jiang R, Du X, Lönnerdal B. Comparison of bioactivities of talactoferrin and lactoferrins from human and bovine milk. *J Pediatr Gastroenterol Nutr* 2014; **59**: 642–52.
  - 113 Yang Z, Jiang R, Chen Q, *et al.* Concentration of Lactoferrin in Human Milk and Its Variation during Lactation in Different Chinese Populations. *Nutrients* 2018; **10**: 1235.
  - 114 Telang S. Lactoferrin: A critical player in neonatal host defense. *Nutrients*. 2018; **10**: 1228.
  - 115 Page MGP. The Role of Iron and Siderophores in Infection, and the Development of Siderophore Antibiotics. *Clin Infect Dis* 2019; **69**: S529–37.
  - 116 Di Lorenzo M, Stork M. Plasmid-Encoded Iron Uptake Systems. *Microbiol Spectr* 2014; **2**: 0030.
  - 117 Bonomo RA. Cefiderocol: A Novel Siderophore Cephalosporin Defeating Carbapenem-resistant Pathogens. *Clin Infect Dis* 2019; **69**: S519–20.
  - 118 Wunderink RG, Wunderink RG, Matsunaga Y, *et al.* LB4. Efficacy and Safety of Cefiderocol vs. High-Dose Meropenem in Patients with Nosocomial Pneumonia—Results of a Phase 3, Randomized, Multicenter, Double-Blind, Non-Inferiority Study. *Open Forum Infect Dis* 2019; **6**: S994–S994.
  - 119 Karlowsky JA, Hackel MA, Tsuji M, Yamano Y, Echols R, Sahm DF. In Vitro Activity of Cefiderocol, a Siderophore Cephalosporin, Against Gram-Negative Bacilli Isolated by Clinical Laboratories in North America and Europe in 2015-2016: SIDERO-WT-2015. *Int J Antimicrob Agents* 2019; **53**: 456–66.
  - 120 Hackel MA, Tsuji M, Yamano Y, Echols R, Karlowsky JA, Sahm DF. In vitro activity of the siderophore cephalosporin, cefiderocol, against carbapenem-nonsusceptible and multidrug-resistant isolates of gram-negative bacilli collected worldwide in 2014 to

2016. *Antimicrob Agents Chemother* 2018; **62**: e01968-17.
- 121 Okomo U, Akpalu NK, Le Doare K, *et al.* Aetiology of invasive bacterial infection and antimicrobial resistance in neonates in sub-Saharan Africa: a systematic review and meta-analysis in line with the STROBE-NI reporting guidelines. *Lancet Infect Dis.* 2019; **19**:1219-1234.
  - 122 Katsarou A, Pantopoulos K. Hepcidin therapeutics. *Pharmaceuticals.* 2018; **11**:127.
  - 123 Ionis Pharmaceuticals. Pipeline. <http://www.ionispharma.com/pipeline/> (accessed Sept 18, 2019).
  - 124 Stefanova D, Raychev A, Arezes J, *et al.* Endogenous hepcidin and its agonist mediate resistance to selected infections by clearing non–transferrin-bound iron. *Blood* 2017; **130**: 245–57.
  - 125 Michels KR, Zhang Z, Bettina AM, *et al.* Hepcidin-mediated iron sequestration protects against bacterial dissemination during pneumonia. *JCI insight* 2017; **2**: e92002.
  - 126 Stefanova D, Raychev A, Deville J, *et al.* Hepcidin Protects against Lethal *Escherichia coli* Sepsis in Mice Inoculated with Isolates from Septic Patients. *Infect Immun* 2018; **86**: e00253-18.
  - 127 Frank KM, Schneewind O, Shieh W-J. Investigation of a Researcher's Death Due to Septicemic Plague. *N Engl J Med* 2011; **364**: 2563–4.
  - 128 Khan FA, Fisher MA, Khakoo RA. Association of hemochromatosis with infectious diseases: expanding spectrum. *Int J Infect Dis* 2007; **11**: 482–7.
  - 129 Bourne, G.; Zhao, L.; Bhandari, A.; Frederick, B.; McMahon, J.; Tran, V.; Annamalai, T.; Mattheakis, L.; Patel, D.; Smythe M. Hepcidin Mimetic PTG-300 for Treatment of Ineffective Erythropoiesis and Chronic Anemia in Hemoglobinopathy Disease. In Proceedings of the 23rd European Hematology Association Congress, Stockholm, Sweden. ; **Abstract #.**
  - 130 Nicholls, A.; Lickliter, J.; Tozzi, L.; Liu, D.; Shames R. Hepcidin mimetic PTG-300 induces dose-related and sustained reductions in serum iron and transferrin

- saturation in healthy subjects. In Proceedings of the 23rd European Hematology Association Congress, Stockholm, Sweden. ; **Abstract #.**
- 131 Lal, A.; Piga, A.; Viprakasit, V.; Maynard, J.; Kattamis, A.; Yaeger, D.; Byrnes, B.; Chawla, L.; Tidmarsh G. A phase 1, open-label study to determine the safety, tolerability, and pharmacokinetics of escalating doses of LJPC-401 (synthetic human hepcidin) in patients with iron overload. In Proceedings of the 23rd European Hematology Association Congress, Stockho. ; **Abstract #.**
  - 132 Aschemeyer S, Qiao B, Stefanova D, *et al.* Structure-function analysis of ferroportin defines the binding site and an alternative mechanism of action of hepcidin. *Blood* 2018; **131**: 899–910.
  - 133 Cai C. Breast Milk Iron Concentrations may be Lower than Previously Reported: Implications for Exclusively Breastfed Infants. *Matern Pediatr Nutr* 2015: **104**:1000104.
  - 134 Frazer DM, Wilkins SJ, Darshan D, Mirciov CSG, Dunn LA, Anderson GJ. Ferroportin Is Essential for Iron Absorption During Suckling, But Is Hyporesponsive to the Regulatory Hormone Hepcidin. *Cell Mol Gastroenterol Hepatol* 2017; **3**: 410–21.
  - 135 Theurl M, Song D, Clark E, *et al.* Mice with hepcidin-resistant ferroportin accumulate iron in the retina. *FASEB J* 2016; **30**: 813–23.
  - 136 Chaston T, Chung B, Mascarenhas M, *et al.* Evidence for differential effects of hepcidin in macrophages and intestinal epithelial cells. *Gut* 2008; **57**: 374–82.
  - 137 Canonne-Hergaux F, Donovan A, Delaby C, Wang HJ, Gros P. Comparative studies of duodenal and macrophage ferroportin proteins. *Am J Physiol - Gastrointest Liver Physiol* 2006; **290**: G156-63.
  - 138 Rao R, Georgieff MK. Iron in fetal and neonatal nutrition. *Semin Fetal Neonatal Med* 2007; **12**: 54–63.

## **ANNEXES**

## ANNEX 1.11 Collaborators and field team details and contributions

Name	Position and Institution	Role in Thesis
Dr Carla Cerami	Senior Investigator Scientist, MRC Unit The Gambia at LSHTM	PhD Primary Supervisor and co-author for manuscripts and papers in Chapters 3-6.
Prof Andrew M. Prentice	Nutrition Theme Leader, MRC Unit The Gambia at LSHTM	Associate Supervisor and co-author for manuscripts and papers in Chapters 3-6.
Prof Hal Drakesmith	Associate Professor of Immunology	Advisory Committee Member.
Prof Simon Andrews	Professor of Molecular Microbiology	Advisory Committee Member.
Dr Sarah Prentice	Lead-Investigator of HYPO-G Study, Keneba Field Station, MRC Unit The Gambia	Lead-Investigator of the pilot study (HYPO-G) to the Neolnnate Study.
Dr Ousman Jarjou	Research Clinician, MRC Unit The Gambia	Research Clinician on the Neolnnate Study and co-author for manuscripts and papers in Chapters 4-6.
Dr Nuradin Ibrahim Mohammed	Study Statistician, MRC Unit The Gambia	Study Statistician on the Neolnnate Study and co-author for manuscripts and papers in Chapters 4-6.
Dr Andrew Armitage	Senior Postdoctoral Scientist	Reviewed the methods of the literature review.
Malcolm Velasco Santiago Rayment Gomez	Intern M.D. and B.Sc. Students, MRC Unit The Gambia	Assistance in study administration and laboratory procedures.
Amulai Touray Edna Anyango Lala Hydara Isatou Dibba	Nutrition Theme Project Coordinator and Assistants, MRC Unit The Gambia	Supported with budget management and study administration.
Tolu Osunnuyi Ebrima Danso Patrick Okot	Laboratory Managers, Keneba Field Station, MRC Unit The Gambia	Laboratory management for Keneba Laboratory were analysis was conducted for Chapters 5 and 6.
Ebrima Sise	Roche Cobas Integra 400+ Technician, Keneba Field Station, MRC Unit The Gambia	Provided technical support for Roche Cobas Integra 400+ seen in Chapter 5 and 6.
Bakary Sonko	Database Manager, Keneba Field Station, MRC Unit The Gambia	Oversaw database design and maintenance for study.
Gilleh Thomas	Database Manager, Fajara, MRC Unit The Gambia	Assistance in database maintenance for study.
Edrisa Sinjanka	Research Nurse Coordinator, Keneba Field Station, MRC Unit The Gambia	Oversaw nurse recruitment and medical training.



Musa Jarjou	Data Entry Manager, Keneba Field Station, MRC Unit The Gambia	Coordination of all data entry activities for study.
Natoma Jarra	Data Entry Clerk, Keneba Field Station, MRC Unit The Gambia	CRF Data entry for study.
Abubacarr Kande	Store Manager, Keneba Field Station, MRC Unit The Gambia	Assistance with local and international laboratory orders to Keneba Field Station.
Ebrima Jallow	Chief Driver, Keneba Field Station, MRC Unit The Gambia	Oversaw transportation of study samples and laboratory materials to Kiang Keneba.
Vivat Thomas-Njie	Coordinator of Clinical Trials Support Office, MRC Unit The Gambia	Advice with compliance with Good clinical Practise and ClinicalTrials.gov.
Haddy Manneh Hulaimatou Zainab Bangura Virginie Jones Morris Ndene Abdou Camara Ebrima Faye James Mendy Sherrifo Darboe	Research SEN Nurses, Fajara, MRC Unit The Gambia	Provided all clinical care to study participants. Conducted all sample and clinical data collection using study protocols.
Sherrifo Jarju Bubacarr JB Touray	Laboratory Technicians, MRC Unit The Gambia	Primary sample processing of all samples during night shifts.
Modou Jeng Tijan Bojang Pa Ousman Nyang Mariama Jallow Omar MS Ceesay	Laboratory Technicians, Kanifing General Hospital	Primary sample processing of all samples during day shifts.
Zaith Babou	Study Driver, MRC Unit The Gambia	Driver for all community visits.
Pa Babou	Laboratory Manager, Kanifing General Hospital	Assistance in the running of the remote laboratory at Kanifing General Hospital.
Kebba Manneh	Chief Executive Officer of Kanifing General Hospital	Supervised all study activities at Kanifing General Hospital.
Haja Zainab Jalloh Binta M Jarju Diminga Mendy Fatoumata Tambi Fatou Cham Fatou Camara Musu Ndura Stella Yawson Maley Camara	Government SEN Nurses and Midwives, Kanifing General Hospital	Assistance in the clinical care of all study participants.
Dr Mass Njie Dr Babading Daffeh	Maternal Ward Clinicians, Kanifing General Hospital	Assistance in the clinical care of all study participants.

Ousainatou Jallow Desmond Ayeh	Procurement Administrators, MRC Unit The Gambia	Assistance in procuring international and local study orders.
Honorary Jammeh	Study Orderly	Oversaw all non-medical study site cleaning and organisation.
Edward Demba	Laboratory Services Manager, MRC Unit The Gambia	Assistance in batch shipment of samples from Fajara to Keneba.
Yaya Giana Lamin Touray	Logistics Administrators, MRC Unit The Gambia	Assistance in importing study material and equipment.
Mariama Balajo	Human Resources Administrator, MRC Unit The Gambia	Assistance in recruitment of study staff.

## ANNEX 1.12 PhD Timeline

PhD Timeline		2015				2016				2017				2018				2019										
		O	N	D	J	F	M	A	M	J	J	A	S	O	N	D	J	F	M	A	M	J	J	A	S	O	N	D
A	PhD Registration and Admin																											
1	Registration																											
2	Upgrading Report Writing																											
3	Upgrading Presentation																											
B	Ethics																											
1	Submission to LSHTM Ethics Committee																											
2	Submission to MRC ING SAM, MRC SCC and MRCG/GG Ethics Committee																											
3	Community Sensitisation																											
4	Open Day																											
C	Study Initiation																											
1	BMGF Grant Writing																											
2	Study Design																											
3	BMGF Funding Approval																											
4	Remote Laboratory Set Up																											
5	Clinical SOP Design																											
6	SSP Design																											
7	Procurement of Study Equipment																											
8	Data Analysis Plan																											
9	Database Design																											
10	Form Pilot Testing																											
11	Study Team Recruitment and Training																											
D	Clinical Section (Kanifing General Hospital - Kombo Area)																											
1	Recruitment of Subjects																											
2	Sample Collection																											
3	Primary Sample Analysis and Processing																											
4	Data Entry																											
5	Clinical Data Cleaning																											
6	Handover of Laboratory, Office and Ward Items (Kanifing General Hospital)																											
E	Laboratory Section (MRC Keneba - Kiang)																											
1	Batch Sample Movement																											
2	Secondary Laboratory Analysis - Roche Cobas Integra 400+																											
3	Secondary Laboratory Analysis - Hepcidin ELISA																											
4	Secondary Laboratory Analysis - G6PD Deficiency Testing																											
F	Data Analysis Section																											
1	Data Analysis																											
2	Archieving Study Forms																											
G	Thesis Preperation																											
1	Methodology/Protocol Paper																											
2	Literature Review																											
3	Main Paper																											
4	Secondary Paper																											
5	Thesis Writing (Introduction, Discussion and Formatting)																											

## ANNEX 1.13 MRCG Scientific Coordinating Committee Ethics Letter



Dr Carla Cerami  
MRC Unit The Gambia  
Keneba

17 February 2017

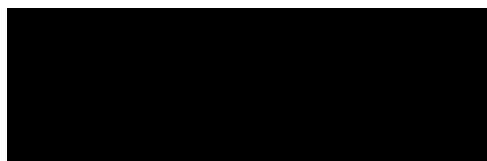
Dear Dr Cerami

### **SCC 1525v1.1, Iron and Infection: Neonatal Nutritional Immunity (Neolnnate Study)**

Thank you for submitting your revised proposal 16 February 2017 addressing the issues raised by the SCC at its meeting held on 6 February 2017.

I have looked at the modified proposal and I am satisfied with the changes you have made according to the SCC queries and recommendations. The proposal can now be forwarded to the Ethics Committee for further consideration at its meeting on 24 February 2017.

With best wishes



Professor Umberto D'Aessandro  
Chair, Scientific Coordinating Committee

#### **Documents submitted for review:**

- Response letter – 14 February 2017
- SCC application form, version 1.1 – 16 February 2017
- Cover letter – 24 January 2017
- ICD, version 1.1 – 15 February 2017
- CVs: Carla Cerami; James Henry Cross
- Budget

#### **Scientific Coordinating Committee**

**MRC Unit The Gambia**

**PO Box 273 Banjul, The Gambia**

**West Africa**

Switchboard (+220) 4495442/6 Ext 2308

Fax (+220) 4495919/4496513

E-mail: [scc@mrc.gm](mailto:scc@mrc.gm)

Intranet: <http://mrcportal/Committees/SCC/SitePages/Home.aspx>

Webpage: <https://mrcportal.mrc.gm/Committees/SCC/SitePages/Home.aspx>

## ANNEX 1.14 The Gambia Government/MRC Joint Ethics Committee Letter

The Gambia Government/MRC Joint

### **ETHICS COMMITTEE**

C/o MRC Unit: The Gambia, Fajara  
P.O. Box 273, Banjul  
The Gambia, West Africa  
Fax: +220 – 4495919 or 4496513  
Tel: +220 – 4495442-6 Ext. 2308  
Email: ethics@mrc.gm

19 April 2017

Dr Carla Cerami  
MRC Unit The Gambia, Keneba

Dear Dr Carla

#### **SCC 1525v1.1, Iron and Infection: Neonatal Nutritional Immunity (Neolnnate Study)**

Thank you for submitting your response letter dated 21 March 2017 addressing the issues raised by The Gambia Government/MRC Joint Ethics Committee at its meeting held on 24 February 2017.

I have also noted the revisions on the informed consent document as per our recommendations. This project has now received full Ethics Committee approval and may proceed.

With best wishes

Yours sincerely



Mr Malamin Sonko  
Chairman, Gambia Government/MRC Joint Ethics Committee

#### **Documents submitted for review:**

- SCC letters – 9 February 2017/17 February 2017
- Response letters – 14 February 2017; 21 March 2017
- SCC application form, version 1.1 – 16 February 2017
- Cover letter – 24 January 2017
- ICD, version 1.2 – 21 March 2017
- CVs: Carla Cerami; James Henry Cross
- Budget

#### **The Gambia Government/MRC Joint Ethics Committee:**

*Mr Malamin Sonko, Chairman  
Prof Ousman Nyan, Scientific Advisor  
Ms Naffie Jobe, Secretary  
Dr Roddie Cole  
Dr Ahmadou Lamin Samateh  
Mrs Tulai Jawara-Ceesay*

*Prof. Umberto D'Alessandro  
Dr Ramatoulie Njie  
Prof Martin Antonio  
Dr Jane Achan  
Dr Momodou L. Waggeh  
Dr Siga Fatima Jagne*

# ANNEX 1.15 London School of Hygiene & Tropical Medicine Ethics Letter

## London School of Hygiene & Tropical Medicine

Keppel Street, London WC1E 7HT  
United Kingdom  
Switchboard: +44 (0)20 7636 8636

[www.lshtm.ac.uk](http://www.lshtm.ac.uk)

LONDON  
SCHOOL of  
HYGIENE  
& TROPICAL  
MEDICINE



### Observational / Interventions Research Ethics Committee

Mr James Cross

LSHTM

22 September 2017

Dear James,

**Study Title:** Iron and Infection: Neonatal Nutritional Immunity (NeoInnate)

**LSHTM Ethics Ref:** 14316

The Observational Committee reviewed the above application.

The documents reviewed were:

Document Type	File Name	Date	Version
Investigator CV	SCCxxxx 22 Jan 2017 (NeoInnate) - Appendix - Co-Investigator J.H.Cross CV	22/01/2017	1
Investigator CV	SCCxxxxx 22Jan 2017 (NeoInnate) - Appendix - PI C.Cerami CV	22/01/2017	1
Protocol / Proposal	SCC1525v1.1 15 Feb 2017 (NeoInnate) - Appendix - Study Design	15/02/2017	1.1
Protocol / Proposal	SCC1525v1.1 15 Feb 2017 (NeoInnate) - Appendix - New Ballard Score Chart	15/02/2017	1.1
Protocol / Proposal	SCC1525v1.1 15 Feb 2017 (NeoInnate) - Appendix - Mock Consort Diagram	15/02/2017	1.1
Protocol / Proposal	SCC1525v1.1 15 Feb 2017 (NeoInnate) - SCC Application Form_CLEAN COPY_APPROVED	15/02/2017	1.1
Information Sheet	SCC1525v1.1 15 Feb 2017 (NeoInnate) - Mother and Neonate - InfoConsent Form CLEAN_APPROVED	15/02/2017	1.1
Local Approval	SCC 1525v1.1_Cerami_GAMBIAN ETHICS Approved_19Apr17	19/04/2017	1.1
Investigator CV	CV_Andrew Prentice	04/07/2017	1

### Request for Clarification

The Committee would like to request further information or clarification before giving their final opinion of the research. The requested further information or clarification is set out below.

The Committee will delegate authority to confirm its final opinion on the application to the Chair.

### Further information or clarification required

- 1) This applicant says they have upgraded, but gives the upgrade date as next month, 20 Sept. Can he clarify?
- 2) Abbreviations. Several non-standard abbreviations were used the application, which were not defined, including 'TSAT' (the primary study outcome) and 'DAG' (relating to the analysis). It would have been helpful if these had been spelled out or more clearly defined.
- 3) Blood draws. Please clarify when, and how frequently, blood samples will be drawn. In addition, please clarify the total amount of blood that will be drawn from pre-term, low birth weight, and full-term neonates over the duration of the study. Please provide a reference for the maximum allowable amount of blood to be drawn safely from neonates, particularly pre-term and low birth weight neonates.
- 4) Information sheet and consent form. The information sheet is extensive and the committee found that it was not well targeted to mothers or parents. Specifically, the number of blood draws and timing of these it is not entirely clear. Although the information sheet states the amount of blood that will be drawn

## ANNEX 1.16 Oral iron acutely elevates bacterial growth in human serum (FeVir Study)

### RESEARCH PAPER COVER SHEET

---

Please note that a cover sheet must be completed for each research paper included within a thesis.

#### SECTION A – Student Details

Student ID Number	LSH158152	Title	Mr.
First Name(s)	James Henry		
Surname/Family Name	Cross		
Thesis Title	Iron and Infection: Neonatal Iron Transition		
Primary Supervisor	Dr Carla Cerami		

If the Research Paper has previously been published please complete Section B, if not please move to Section C.

#### SECTION B – Paper already published

Where was the work published?	Nature Scientific Reports			
When was the work published?	23 <sup>rd</sup> November 2015			
If the work was published prior to registration for your research degree, give a brief rationale for its inclusion	It was published during my registration and has been included as to provide further evidence that high levels of serum iron support the bacterial growth.			
Have you retained the copyright for the work?*	<table><tbody><tr><td>Yes</td><td>Was the work subject to academic peer review?</td><td>Yes</td></tr></tbody></table>	Yes	Was the work subject to academic peer review?	Yes
Yes	Was the work subject to academic peer review?	Yes		

\*If yes, please attach evidence of retention. If no, or if the work is being included in its published format, please attach evidence of permission from the copyright holder (publisher or other author) to include this work.

#### **Author Contributions**

J.H.C., R.S.B., A.P. and C.C. designed the experiments. J.H.C. and A.T.J. performed experiments. J.H.C., R.S.B. and R.W. recruited patients; A.F., A.P. and C.C. analyzed data; A.P. and C.C. wrote the manuscript. All authors approved the final manuscript.

#### **Additional Information**

**Competing financial interests:** The authors declare no competing financial interests.

**How to cite this article:** Cross, J. H. *et al.* Oral iron acutely elevates bacterial growth in human serum. *Sci. Rep.* 5, 16670; doi: 10.1038/srep16670 (2015).



This work is licensed under a Creative Commons Attribution 4.0 International License. The images or other third party material in this article are included in the article's Creative Commons license, unless indicated otherwise in the credit line; if the material is not included under the Creative Commons license, users will need to obtain permission from the license holder to reproduce the material. To view a copy of this license, visit <http://creativecommons.org/licenses/by/4.0/>

### **SECTION C – Prepared for publication, but not yet published**

Where is the work intended to be published?

Please list the paper's authors in the intended authorship order:

Stage of publication

Choose an item.

### **SECTION D – Multi-authored work**

For multi-authored work, give full details of your role in the research included in the paper and in the preparation of the paper. (Attach a further sheet if necessary)

James H. Cross designed and performed the experiments. He also recruited patients and approved the final manuscript.

### **SECTION E**

**Student Signature**



**Date**

08/11/2019

**Supervisor Signature**



**Date**

08/11/2019



# SCIENTIFIC REPORTS

OPEN

## Oral iron acutely elevates bacterial growth in human serum

James H. Cross<sup>1</sup>, Richard S. Bradbury<sup>1,2</sup>, Anthony J. Fulford<sup>1,3</sup>, Amadou T. Jallow<sup>1</sup>, Rita Wegmüller<sup>1</sup>, Andrew M. Prentice<sup>1,3</sup> & Carla Cerami<sup>4</sup>

Received: 30 May 2015

Accepted: 01 October 2015

Published: 23 November 2015

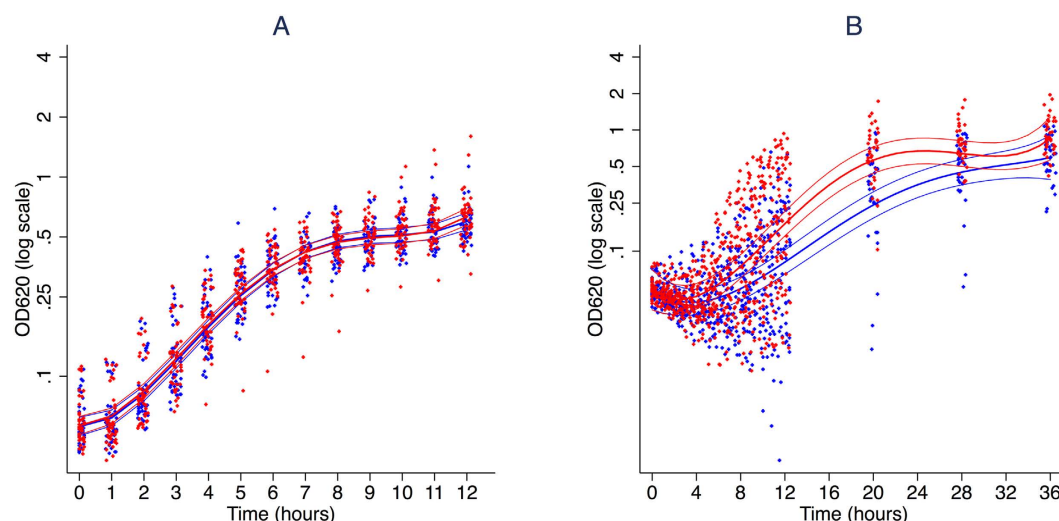
Iron deficiency is the most common nutrient deficiency worldwide and routine supplementation is standard policy for pregnant mothers and children in most low-income countries. However, iron lies at the center of host-pathogen competition for nutritional resources and recent trials of iron administration in African and Asian children have resulted in significant excesses of serious adverse events including hospitalizations and deaths. Increased rates of malaria, respiratory infections, severe diarrhea and febrile illnesses of unknown origin have all been reported, but the mechanisms are unclear. We here investigated the *ex vivo* growth characteristics of exemplar sentinel bacteria in adult sera collected before and 4 h after oral supplementation with 2 mg/kg iron as ferrous sulfate. *Escherichia coli*, *Yersinia enterocolitica* and *Salmonella enterica* serovar Typhimurium (all gram-negative bacteria) and *Staphylococcus epidermidis* (gram-positive) showed markedly elevated growth in serum collected after iron supplementation. Growth rates were very strongly correlated with transferrin saturation ( $p < 0.0001$  in all cases). Growth of *Staphylococcus aureus*, which preferentially scavenges heme iron, was unaffected. These data suggest that even modest oral supplements with highly soluble (non-physiological) iron, as typically used in low-income settings, could promote bacteremia by accelerating early phase bacterial growth prior to the induction of immune defenses.

Iron deficiency (ID) remains the most pervasive nutritional deficiency worldwide. The prevalence of ID in mothers and young children frequently exceeds 50% in low-income countries. Insufficient iron impairs growth and cognitive development in childhood<sup>1,2</sup>.

Low cost iron supplements are effective for the treatment of ID and in countries with ID rates of >40%, the World Health Organization recommends universal iron supplementation of pregnant women and young children<sup>3–5</sup>. To overcome perceived limitations in the ability to absorb iron, supplements usually employ highly soluble forms of iron (ferrous sulfate or fumarate) given in rather large non-physiological bolus doses. The wisdom of these policies has long been questioned<sup>6,7</sup> and has come under serious scrutiny starting in 2006 with the premature termination of a large trial in Pemba, Tanzania after significant increases in serious adverse outcomes (hospitalizations and deaths) in young children receiving iron-folate supplements were seen<sup>8</sup>. The emphasis was originally focused on malaria as the causative agent for the increases in morbidity and mortality during iron supplementation<sup>9</sup> but subsequent trials have described excesses of other infections in groups randomized to iron or multiple micronutrients containing iron<sup>10–15</sup>. These findings have paralyzed iron supplementation policies.

The underlying mechanisms and the types of organisms responsible for these clinical and epidemiological observations remain unclear. Recent field studies suggest that oral iron supplementation in children increases susceptibility to bacterial infections, particularly diarrhea<sup>13,16</sup>, alters the gut microbiota<sup>16,17</sup>, and increases the virulence of many common bacterial enteropathogens<sup>18–20</sup>.

<sup>1</sup>MRC Keneba, MRC Unit, The Gambia, Atlantic Blvd, Serrekunda, Gambia. <sup>2</sup>School of Medical and Applied Sciences, Central Queensland University, North Rockhampton, Queensland, Australia. <sup>3</sup>MRC International Nutrition Group, London School of Hygiene & Tropical Medicine, Keppel Street, WC1E 7HT, London, UK. <sup>4</sup>Department of Epidemiology, Gillings' School of Global Public Health, University of North Carolina, Chapel Hill, North Carolina, USA. Correspondence and requests for materials should be addressed to C.C. (email: ccerami@unc.edu)



**Figure 1. Growth of sentinel gram-positive bacteria in human serum before and after oral iron supplementation.** *S. aureus* (A) and *S. epidermidis* (B) were grown in serum from subjects before (blue) and after (red) oral iron supplementation with 400 mg ferrous sulfate (containing the equivalent of 130 mg of elemental iron). The thicker central lines represent the fitted curves for an average individual and the thinner lines the 95% confidence intervals for these estimates. The points show individual values of OD<sub>620</sub>. Curves were derived from mixed effects models fitting degree-four orthogonal polynomials in time to log(OD<sub>620</sub>), where OD<sub>620</sub> is the mean optical density of the three replicates at time (t).

Numerous animal studies over many decades have shown that administration of iron in diverse forms accelerates the growth of peritoneally-injected pathogens, causing a septicemia with rapidly fatal outcomes<sup>21,22</sup>. We here examine the possibility in humans that a simple oral dose of supplemental iron could promote bacterial growth in serum. We used a series of *ex vivo* bacterial growth assays with sentinel organisms that were selected on the basis of their varying modes of pathogenesis and abilities to scavenge iron from the host.

## Results

**Oral iron supplementation increases iron parameters in serum.** To determine the effects of oral iron supplementation on bacterial growth in human serum, we enrolled 48 normal healthy non-anemic male subjects [mean  $\pm$  SD: Hemoglobin (Hgb) =  $14.5 \pm 1.13$  g/dL; Mean Corpuscular Volume (MCV) =  $83.8 \pm 5.5$  fL; Ferritin =  $62.8 \pm 53.2$  ng/mL]. Volunteers donated serum immediately before, and then four hours after, oral ingestion of 400 mg ferrous sulfate (containing the equivalent of 130 mg of elemental iron). Transferrin saturation (TSAT) increased from 42.1% ( $\pm 12.5\%$ , SD) to 75.7% ( $\pm 18.1\%$ , SD) and total serum iron increased from 30.3  $\mu$ mol/L ( $\pm 10.2 \mu$ mol/L, SD) to 53.0  $\mu$ mol/L ( $\pm 15.8 \mu$ mol/L, SD) four hours after iron supplementation.

**Effects of oral iron supplementation on *ex vivo* bacterial growth in serum.** We next measured the growth of the five species of sentinel bacteria in the baseline and post-dose sera. To account for the between-subject variance in the starting transferrin saturation levels, we used mixed statistical models to allow two nested higher levels of variation: patient and bleed (pre- and post-iron supplementation). This enabled us to independently analyze the effects of TSAT and iron supplementation.

*S. aureus*, an organism with a strong preference for heme-derived iron<sup>23</sup>, behaved differently from the other four bacteria (Fig. 1A). Iron supplementation had no impact on the general pattern of the growth curve ( $p = 0.3$ ). Both the time to reach peak doubling time ( $p = 0.21$ ), and the doubling time during the exponential growth phase ( $p = 0.78$ ) were also unchanged by iron supplementation (Table 1). Growth did not correlate with TSAT ( $p = 0.08$ ) (Table 2).

*S. epidermidis* (Fig. 1B) demonstrated an initial delay in growth in comparison with the other species. Iron supplementation influenced the overall pattern of the growth curve ( $p < 0.0001$ ). Specifically, iron supplementation reduced the lag phase, the time to reach peak doubling time ( $p = 0.001$ ), and increased the doubling time during the exponential growth phase ( $p < 0.001$ ) (Table 1). The very strong effect of iron on the overall increase in bacterial growth ( $X^2 = 55$  (approximately),  $p < 0.0001$ ) was equally explained using pre/post supplementation as a dichotomized variable or by TSAT (Table 2).

*S. Typhimurium* (Fig. 2A) and *E. coli* (Fig. 2B) both showed highly significant differences in their growth curves after iron supplementation ( $p < 0.0001$ ). The doubling times post-iron supplementation were significantly shorter than pre-iron supplementation ( $p < 0.0001$ ), but time to reach peak doubling time was unaffected by treatment (Table 1). Transferrin saturation had a very strong effect on overall

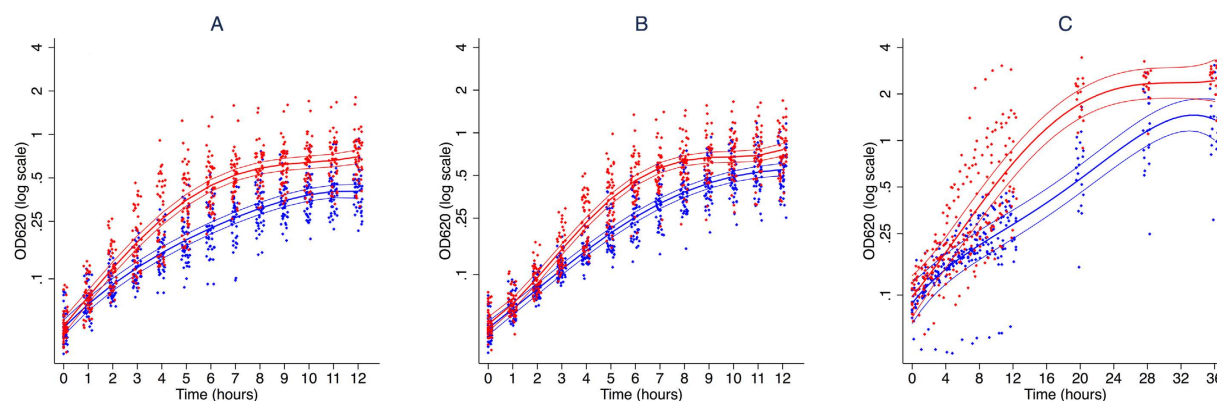
Analysis of Growth		<i>S. aureus</i>	<i>S. epidermidis</i>	<i>E. coli</i>	<i>S. Typhimurium</i>	<i>Y. enterocolitica</i>
Growth Curve	X <sup>2</sup> (5 df)	6.05	65.70	225.00	232.00	117.70
	p-value	0.30	<0.0001	<0.0001	<0.0001	<0.0001
Time to Reach Peak Doubling Time (hours)	Mean pre (CI95%)	3.34 (3.21, 3.47)	13.7 (11.8, 15.7)	1.86 (0.50, 3.21)	Not Applicable - No MAX	Not Applicable - No MAX
	Mean post (CI95%)	3.25 (3.12, 3.38)	10.70 (10.0, 11.4)	2.56 (2.32, 2.79)		
	z	−1.26	−3.24	1.03		
	p-value	0.21	0.001	0.3		
Doubling Time During Exponential Phase (hours)	Mean pre (CI95%)	1.74 (1.67, 1.81)	4.82 (4.2, 5.5)	2.14 (1.99, 2.29)	2.03 (1.81, 2.26)	4.96 (3.0, 7.0)
	Mean post (CI95%)	1.75 (1.68, 1.82)	3.36 (3.0, 3.7)	1.50 (1.44, 1.56)	1.60 (1.37, 1.62)	4.37 (2.8, 6.0)
	z	0.28	−4.56	−8.04	−4.34	−1.73
	p-value	0.78	<0.001	<0.0001	<0.001	0.62

**Table 1. Statistical analysis of *ex vivo* bacterial growth assays.** For each species of bacterium we compared its growth pre- and post-iron supplementation by examining differences in (1) Growth Curve, i.e. general pattern curve of the overall growth trajectories; (2) Time to Reach Peak Doubling Time, i.e. the time at which the rate of increase in log(OD<sub>620</sub>) was at a maximum; and (3) Doubling Time During Exponential Growth Phase.

Hypothesis Testing		<i>S. aureus</i>	<i>S. epidermidis</i>	<i>E. coli</i>	<i>S. Typhimurium</i>	<i>Y. enterocolitica</i>
(1) Iron supp. affects growth	X <sup>2</sup> (5 df)	5.92	55.70	221.00	213.00	108.17
	p-value	0.31	<0.0001	<0.0001	<0.0001	<0.0001
(2) Iron supp. affects growth independently of TSAT	X <sup>2</sup> (5 df)	8.85	7.53	35.10	22.40	32.71
	p-value	0.12	0.18	<0.0001	0.0004	<0.0001
(3) TSAT affects growth	X <sup>2</sup> (5 df)	9.79	55.50	300.00	348.00	120.41
	p-value	0.08	<0.0001	<0.0001	<0.0001	<0.0001
(4) TSAT affects growth independently of iron supplementation	X <sup>2</sup> (5 df)	12.70	6.22	69.50	105.00	36.89
	p-value	0.03	0.29	<0.0001	<0.0001	<0.0001

**Table 2. Statistical testing of the independent effects of iron supplementation and transferrin saturation (TSAT).** Three models (mathematical equations) were fitted for each bacterial species: (a) iron supplementation × (t1 t2 t3 t4), i.e. iron supplementation, the time polynomials and their interactions; (b) TSAT × (t1 t2 t3 t4), i.e. TSAT, the time polynomials and their interactions; (c) (iron supplementation + TSAT) × (t1 t2 t3 t4), i.e. iron supplementation, TSAT and both their interactions with the time polynomials. We employed the likelihood ratio test to compare models for which the growth patterns were and were not dependent on TSAT or iron supplementation. The effects of iron supplementation and TSAT were derived from models (a) and (b) respectively and refer to the joint effects of the variable and the terms for its interaction with the time polynomials. The independent (conditional) effects of iron supplementation after controlling for TSAT, and TSAT after controlling for iron supplementation, are both derived from model (c) and again refer to the joint effects of the variable and the terms for its interaction with the time polynomials. For each bacterial species, each of the following four hypotheses were tested: (1) iron supplementation has an impact on bacterial growth; (2) iron supplementation has an impact on bacterial growth independently of TSAT; (3) TSAT has an impact on bacterial growth; and (4) TSAT has an impact on bacterial growth independently of iron supplementation. Significance was determined using a chi-squared test. Chi-squared and p-values are reported.

growth rates of both *S. Typhimurium* ( $X^2 = 348$ ,  $p < 0.0001$ ) and *E. coli* ( $X^2 = 300$ ,  $p < 0.0001$ ), comparable to, indeed a little larger than, that of iron supplementation in both *S. Typhimurium* ( $X^2 = 213$ ,  $p < 0.0001$ ) and *E. coli* ( $X^2 = 221$ ,  $p < 0.0001$ ). For *E. coli*, iron supplementation ( $X^2 = 35$ ,  $p < 0.0001$ ) and TSAT ( $X^2 = 69$ ,  $p < 0.0001$ ) each had significant effects on bacterial growth after controlling for the other. The same held true for *S. Typhimurium* growth where both iron supplementation ( $X^2 = 22$ ,  $p = 0.0004$ ) and TSAT ( $X^2 = 105$ ,  $p < 0.0001$ ) each had significant effects on bacterial growth after controlling for the other (Table 2). This is likely to be because, in addition to capturing the effect of supplementation, transferrin saturation also explains differences between individuals.



**Figure 2. Growth of sentinel gram-negative bacteria in human serum before and after oral iron supplementation.** *S. Typhimurium* (A), *E. coli* (B) and *Y. enterocolitica* (C) were grown in serum from subjects before (blue) and after (red) oral iron supplementation with 400 mg ferrous sulfate (containing the equivalent of 130 mg of elemental iron). The thicker central lines represent the fitted curves for an average individual and the thinner lines the 95% confidence intervals for these estimates. The points show individual values of OD<sub>620</sub>. Curves were derived from mixed effects models fitting degree-four orthogonal polynomials in time to log(OD<sub>620</sub>), where OD<sub>620</sub> is the mean optical density of the three replicates at time (t).

For *Y. enterocolitica* (Fig. 2C), the maximum growth rate was difficult to locate. In fact, although the growth curves clearly differ significantly ( $p < 0.0001$ ), the doubling times at one hour did not differ significantly. As was the case for *S. Typhimurium* and *E. coli*, TSAT had an impact on growth ( $X^2 = 120$ ,  $p < 0.0001$ ) as did iron supplementation ( $X^2 = 108$ ,  $p < 0.0001$ ). Additionally, iron supplementation ( $X^2 = 33$ ,  $p < 0.0001$ ) and TSAT ( $X^2 = 37$ ,  $p < 0.0001$ ) each had significant effects on bacterial growth after controlling for the other (Table 2).

## Discussion

The biologically useful redox characteristics of the Fe(II) to Fe(III) transition place iron apart from other nutrients. Additionally, it lies at the epicenter of the host-pathogen battle for resource control. Host defense mechanisms to withhold iron from invading pathogens are some of the most evolutionarily conserved innate strategies against infection<sup>24</sup>, but most bacterial species have evolved counter-acting strategies for pirating host iron including: (1) receptors that bind transferrin, lactoferrin or hemoglobin; and (2) low molecular weight siderophores that acquire iron from host proteins or from low molecular weight iron compounds<sup>25</sup>.

The potential health threat posed by exogenous iron, repeatedly demonstrated in animal models<sup>25</sup>, has tended to be overlooked in clinical settings. The recent iron trials with adverse outcomes in children in developing countries<sup>8,10–15</sup> have prompted new mechanistic studies providing experimental verification that oral iron adversely modifies the gut microbiome<sup>17,26</sup> and increases the virulence of pathogenic enteric bacteria<sup>18,20</sup>. In this study we focused on the issue of systemic, as opposed to enteric, bacterial infections building upon prior knowledge that iron can precipitate septicemias (for instance, based on the disastrous outcomes of intramuscular iron-dextran administration to Polynesian neonates<sup>27</sup>).

The *ex vivo* assays we describe here show that customary oral supplementation with highly-soluble iron as ferrous sulfate can profoundly affect the growth dynamics of four of the five sentinel species we studied. This could potentially undermine a key component of innate immunity allowing such organisms to achieve overwhelming numbers by the time adaptive immune defense mechanisms are up-regulated. Note that the very strong correlations between TSAT and growth rates emphasizes the importance of this variable even in the presence of likely inter-individual differences in other iron-related (e.g. lipocalin-2, haptoglobin) and other (e.g. defensins) non-cellular defense mechanisms within the sera.

Recent molecular insights into human iron metabolism have challenged the basic pillars on which public health strategies involving highly soluble iron supplements have been developed. The prior belief that humans are constitutionally inefficient at absorbing iron, and hence require large non-physiological doses taken apart from food, is now overturned by the knowledge that hepcidin actively down-regulates iron acquisition especially in the presence of an infectious threat<sup>24</sup>. The dual regulation of hepcidin by iron and infection (inflammation) underscores the threat posed by exogenous iron. An increase of hepcidin caused by an infection might have the evolutionary function of decreasing further iron uptake from the intestine to reduce circulating iron fuel for microorganisms. This suggests that we should not interfere via high dose iron supplements.

Transferrin saturation is homeostatically controlled with a normal range between 15–50% in males. Our data show that increasing TSAT from a mean of 42% to 76% profoundly stimulated bacterial growth with a continuous association across all levels of TSAT. The role of TSAT in mediating host susceptibility to infection has been known for almost half a century<sup>28</sup>. However, neither the strength of this association, nor the ability of acute increases in TSAT following iron doses to so rapidly favor bacterial growth, have been previously appreciated. The dose level selected for the adults in this study was based on that most frequently used for young children with iron deficiency (2 mg/kg/day), however our subjects were iron replete. In iron deficient children, hepcidin would be down-regulated to allow maximal iron absorption<sup>29</sup> and hence TSAT would be expected to rise even further. Although a higher percentage of increase can be expected, the TSAT baseline will be lower and may still end up lower than in iron replete men. In the event of accidental ingress of pathogens through a cut, abrasion or leaky gut, these high levels of TSAT could precipitate a fulminant bacteremia before other cognate immune defenses have time to respond.

Iron absorbed from a natural food matrix, or even when ferrous sulfate is given with food<sup>30</sup>, is released much more slowly and causes minor deviations in TSAT and in non-transferrin bound iron (NTBI)<sup>30</sup> which, according to the data presented here, would be much less likely to promote bacterial growth. In regions of the world where iron-rich foods are scarce, or too costly, supplementation with slow release nano-molecular formulations that mimic a food matrix<sup>31</sup> may provide a safer option and might additionally reduce the adverse effects of unabsorbed iron on the gut microbiota. Field trials of such compounds are warranted. Additional safe approaches to supplement iron include the use of oral bovine lactoferrin<sup>32</sup>.

## Materials and Methods

**Subjects.** Forty-eight male Gambian subjects (averaging 40y; range 21–64y) were recruited. Subjects had no history of fever, illness or anti-microbial use during the preceding seven days; were malaria rapid test negative and were non-anemic (Hemoglobin >12 g/dL). All patients donated blood between 9 and 10 am, on an empty stomach, immediately prior to and four hours after taking 400 mg ferrous sulfate orally. Food was provided two hours after iron supplementation.

Informed consent was obtained from all subjects. This study was approved by the Gambian Government/MRC Joint Ethics Committee (SCC1312v2) and by the University of North Carolina Institutional Review Board (protocol #143044). All experiments were performed in accordance with the approved guidelines.

**Biochemical Parameters.** Complete blood counts were obtained using a Medonic M series (Boule Diagnostics Int AB, Stockholm, Sweden) hematology analyzer. Serum biochemical parameters including serum iron, transferrin, ferritin and transferrin saturation were obtained using a Cobas Integra 400 plus (Roche, Basel, Switzerland) biochemistry analyzer.

**Bacterial Growth assays.** *Staphylococcus aureus* (strain NCTC8325), *Staphylococcus epidermidis* (FDA strain PCI1200, ATCC12228), *Salmonella enterica* serovar Typhimurium (strain LT2, ATCC19585) and *Escherichia coli* (strain Crooks, ATCC8739) were grown overnight for 18 hours at 37 °C in 5 mL iron free minimal growth media, Iscove's Modified Dulbecco's Medium (IMDM, Invitrogen). This was conducted in air with continuous shaking (250 rpm). A high-virulence, siderophore producing *Yersinia enterocolitica* (strain WA-314, ATCC51871) was grown in IMDM containing 10 mM ethylene glycol tetraacetic acid (EGTA, Sigma) (pH7). All growth assays were run in triplicate in IMDM containing 50% heat-inactivated human serum. Bacterial growth was monitored by measuring the optical density at 620 nm (OD<sub>620</sub>) hourly for 12 hours (*Staphylococcus aureus*, *Salmonella enterica* serovar Typhimurium, and *Escherichia coli*) and then at 20, 28, 36 hours (*Staphylococcus epidermidis* and *Yersinia enterocolitica*) using a Multiscan FC ELISA plate reader (Thermo Scientific).

**Statistical analysis.** We analyzed the growth assays with a mixed model using a quartic polynomial in Stata12 (StataCorp, College Station, TX). We used the logarithm of the mean of the three replicate OD readings at each time point as the response variable and modeled growth trajectories by fitting this to orthogonal polynomials (to degree 4) in time using mixed effects models with random intercept and coefficients due to patient and, nested within patient, bleed (pre and post). In effect, therefore, the growth trajectory for the  $i^{\text{th}}$  blood sample was modeled by an equation of the form:

$$\log[\text{OD}_i(t)] = (\beta_0 + \tau_{0i}) + (\beta_1 + \tau_{1i})t + (\beta_2 + \tau_{2i})t^2 + (\beta_3 + \tau_{3i})t^3 + (\beta_4 + \tau_{4i})t^4 + \varepsilon_{it},$$

where  $t$  is time since inoculation, the  $\beta$ s are estimated coefficients and the  $\tau$ s and  $\varepsilon$  are the random effects.

For each species of bacterium, we compared its growth pre- and post-iron supplementation by examining differences in (1) Growth Curve i.e. general pattern of the overall growth trajectories; (2) Time to Reach Peak Doubling Time, i.e. the time at which the rate of increase in  $\log[\text{OD}]$  was at a maximum and (iii) Doubling Time During Exponential Growth Phase, i.e. when the doubling time was at its greatest, or, if the lag phase was not detectable at 1 hour after inoculation. We estimated the timing of the maximum slope by setting the second derivative with respect to time of the deterministic component of the model to zero. We calculated the doubling time at time  $t$  as  $\log(2)/(\text{slope of } \log[\text{OD}(t)])$ . We employed



the likelihood ratio test to compare models for which the growth patterns were and were not dependent on the transferrin saturation or iron supplementation. We used the delta method to obtain the 95% confidence intervals for the time at which maximum slope occurred and the doubling time.

## References

- Black, M. M., Quigg, A. M., Hurley, K. M. & Pepper, M. R. Iron deficiency and iron-deficiency anemia in the first two years of life: strategies to prevent loss of developmental potential. *Nutr. Rev.* **69** Suppl 1, S64–70 (2011).
- Miller, J. L. Iron deficiency anemia: a common and curable disease. *Cold Spring Harb. Perspect. Med.* **3**, (2013).
- Peña-Rosas, J. P., De-Regil, L. M., Dowsell, T. & Viteri, F. E. Daily oral iron supplementation during pregnancy. *Cochrane Database Syst. Rev.* **12**, (2012) CD004736.
- Okebe, J. U., Yahav, D., Shbita, R. & Paul, M. Oral iron supplements for children in malaria-endemic areas. *Cochrane Database Syst. Rev. Online* CD006589 (2011), doi: 10.1002/14651858.CD006589.pub3.
- Nutritional anaemias. Report of a WHO scientific group. *World Health Organ. Tech. Rep. Ser.* **405**, 5–37 (1968).
- Murray, M. J., Murray, A. B., Murray, M. B. & Murray, C. J. The adverse effect of iron repletion on the course of certain infections. *Br. Med. J.* **2**, 1113–1115 (1978).
- Oppenheimer, S. J. Iron and its relation to immunity and infectious disease. *J. Nutr.* **131**, 616S–633S (2001).
- Sazawal, S. *et al.* Effects of routine prophylactic supplementation with iron and folic acid on admission to hospital and mortality in preschool children in a high malaria transmission setting: community-based, randomised, placebo-controlled trial. *Lancet* **367**, 133–143 (2006).
- Spottiswoode, N., Duffy, P. E. & Drakesmith, H. Iron, anemia and hepcidin in malaria. *Front. Pharmacol.* **5**, 125 (2014).
- Esan, M. O. *et al.* Iron supplementation in HIV-infected Malawian children with anemia: a double-blind, randomized controlled trial. *Clin. Infect. Dis. Off. Publ. Infect. Dis. Soc. Am.* (2013), doi: 10.1093/cid/cit528.
- Jonker, F. A. M. *et al.* Iron status predicts malaria risk in Malawian preschool children. *PLoS One* **7**, e42670 (2012).
- Manno, D. *et al.* Effect of multiple micronutrient-fortified food on mild morbidity and clinical symptoms in Zambian infants: results from a randomised controlled trial. *Eur. J. Clin. Nutr.* **65**, 1163–1166 (2011).
- Soofi, S. *et al.* Effect of provision of daily zinc and iron with several micronutrients on growth and morbidity among young children in Pakistan: a cluster-randomised trial. *Lancet* **382**, 29–40 (2013).
- Veenemans, J. *et al.* Effect of supplementation with zinc and other micronutrients on malaria in Tanzanian children: a randomised trial. *PLoS Med.* **8**, (2011) e1001125.
- Zlotkin S. *et al.* Effect of iron fortification on malaria incidence in infants and young children in Ghana: A randomized trial. *JAMA* **310**, 938–947 (2013).
- Jaeggi, T. *et al.* Iron fortification adversely affects the gut microbiome, increases pathogen abundance and induces intestinal inflammation in Kenyan infants. *Gut* (2014), doi: 10.1136/gutjnl-2014-307720.
- Zimmermann, M. B. *et al.* The effects of iron fortification on the gut microbiota in African children: a randomized controlled trial in Cote d'Ivoire. *Am. J. Clin. Nutr.* **92**, 1406–1415 (2010).
- Kortman, G. A. M., Boleij, A., Swinkels, D. W. & Tjalsma, H. Iron availability increases the pathogenic potential of *Salmonella typhimurium* and other enteric pathogens at the intestinal epithelial interface. *PLoS One* **7**, (2012) e29968.
- Kortman, G. A. *et al.* Low dietary iron intake restrains the intestinal inflammatory response and pathology of enteric infection by food-borne bacterial pathogens. *Eur. J. Immunol.* (2015), doi: 10.1002/eji.201545642.
- Dostal, A. *et al.* *Salmonella* adhesion, invasion and cellular immune responses are differentially affected by iron concentrations in a combined *in vitro* gut fermentation-cell model. *PLoS One* **9**, e93549 (2014).
- Bullen, J. J., Leigh, L. C. & Rogers, H. J. The effect of iron compounds on the virulence of *Escherichia coli* for guinea-pigs. *Immunology* **15**, 581–588 (1968).
- Bullen, J. J., Rogers, H. J. & Griffiths, E. Role of iron in bacterial infection. *Curr. Top. Microbiol. Immunol.* **80**, 1–35 (1978).
- Skaar, E. P., Humayun, M., Bae, T., DeBord, K. L. & Schneewind, O. Iron-source preference of *Staphylococcus aureus* infections. *Science* **305**, 1626–1628 (2004).
- Drakesmith, H. & Prentice, A. M. Hepcidin and the iron-infection axis. *Science* **338**, 768–772 (2012).
- Weinberg, E. D. Iron availability and infection. *Biochim. Biophys. Acta* **1790**, 600–605 (2009).
- Dostal, A. *et al.* Effects of iron supplementation on dominant bacterial groups in the gut, faecal SCFA and gut inflammation: a randomised, placebo-controlled intervention trial in South African children. *Br. J. Nutr.* **112**, 547–556 (2014).
- Barry, D. M. & Reeve, A. W. Increased incidence of gram-negative neonatal sepsis with intramuscular iron administration. *Pediatrics* **60**, 908–912 (1977).
- Kochan, I. Mechanism of tuberculostasis in mammalian serum. I. Role of transferrin in human serum tuberculostasis. *J. Infect. Dis.* **119**, 11–18 (1969).
- Prentice, A. M. *et al.* Hepcidin is the major predictor of erythrocyte iron incorporation in anemic African children. *Blood* **119**, 1922–1928 (2012).
- Schümann, K. *et al.* Impact of oral iron challenges on circulating non-transferrin-bound iron in healthy Guatemalan males. *Ann. Nutr. Metab.* **60**, 98–107 (2012).
- Pereira, D. I. A. *et al.* Nanoparticulate iron(III) oxo-hydroxide delivers safe iron that is well absorbed and utilised in humans. *Nanomedicine Nanotechnol. Biol. Med.* **10**, 1877–1886 (2014).
- Nappi, C. *et al.* Efficacy and tolerability of oral bovine lactoferrin compared to ferrous sulfate in pregnant women with iron deficiency anemia: a prospective controlled randomized study. *Acta Obstet. Gynecol. Scand.* **88**, 1031–1035 (2009).

## Acknowledgements

The authors would like to acknowledge Bakary Darboe, Pierre Coulin, Karamo Camara, Ebrima Sise, Morgan Goheen, Daniel Gardner, Abdoulie Faal and all of the lab staff at the MRC-ING Keneba lab for their endless help, support and enthusiasm during this project. In addition, we would like to thank all of the blood donors who participated in the study. Funding: This work was supported by MCA760-5QX00 to the MRC International Nutrition Group by the UK Medical Research Council (MRC) and the UK Department for International Development (DFID) under the MRC/DFID Concordat agreement, and by the National Institutes of Health and the National Institute of Child Health and Human Development (U01 HD061234 05S2 to CC).

### Author Contributions

J.H.C., R.S.B., A.P. and C.C. designed the experiments. J.H.C. and A.T.J. performed experiments. J.H.C., R.S.B. and R.W. recruited patients; A.F., A.P. and C.C. analyzed data; A.P. and C.C. wrote the manuscript. All authors approved the final manuscript.

### Additional Information

**Competing financial interests:** The authors declare no competing financial interests.

**How to cite this article:** Cross, J. H. *et al.* Oral iron acutely elevates bacterial growth in human serum. *Sci. Rep.* **5**, 16670; doi: 10.1038/srep16670 (2015).



This work is licensed under a Creative Commons Attribution 4.0 International License. The images or other third party material in this article are included in the article's Creative Commons license, unless indicated otherwise in the credit line; if the material is not included under the Creative Commons license, users will need to obtain permission from the license holder to reproduce the material. To view a copy of this license, visit <http://creativecommons.org/licenses/by/4.0/>

## ANNEX 3.8 New Ballard Score Sheet (Gestational Aging)

### The New Ballard Score

www.ballardscore.com

#### NEUROMUSCULAR MATURITY

SIGN	SCORE							SIGN SCORE
	-1	0	1	2	3	4	5	
Posture								
Square Window								
Arm Recoil								
Popliteal Angle								
Scarf Sign								
Heel To Ear								
TOTAL NEUROMUSCULAR SCORE								

#### MATURITY RATING

TOTAL SCORE	WEEKS
-10	20
-5	22
0	24
5	26
10	28
15	30
20	32
25	34
30	36
35	38
40	40
45	42
50	44

SIGN	SCORE							SIGN SCORE
	-1	0	1	2	3	4	5	
Skin	Sticky, friable, transparent	gelatinous, red, translucent	smooth pink, visible veins	superficial peeling &/or rash, few veins	cracking, pale areas, rare veins	parchment, deep cracking, no vessels	leathery, cracked, wrinkled	
Lanugo	none	sparse	abundant	thinning	bald areas	mostly bald		
Plantar Surface	heel-toe 40-50mm: -1 <40mm: -2	>50 mm no crease	faint red marks	anterior transverse crease only	creases ant. 2/3	creases over entire sole		
Breast	imperceptible	barely perceptible	flat areola no bud	stippled areola 1-2 mm bud	raised areola 3-4 mm bud	full areola 5-10 mm bud		
Eye / Ear	lids fused loosely: -1 tightly: -2	lids open pinna flat stays folded	sl. curved pinna; soft; slow recoil	well-curved pinna; soft but ready recoil	formed & firm instant recoil	thick cartilage ear stiff		
Genitals (Male)	scrotum flat, smooth	scrotum empty, faint rugae	testes in upper canal, rare rugae	testes descending, few rugae	testes down, good rugae	testes pendulous, deep rugae		
Genitals (Female)	clitoris prominent & labia flat	prominent clitoris & small labia minora	prominent clitoris & enlarging minora	majora & minora equally prominent	majora large, minora small	majora cover clitoris & minora		
TOTAL PHYSICAL MATURITY SCORE								

#### Gestation by Dates

	weeks
--	-------

Birth date	Hour	am	pm

APGAR	1 min	5min

#### Scoring

Gest. Age by Maturity Rating	_____ weeks
Time of Exam	Date _____ am Hour _____ pm
Age at Exam	_____ hours

Signature of Examiner

M.D. / R.N.

#### References :

Ballard JL, Khoury JC, Wedig K, et al: New Ballard Score, expanded to include extremely premature infants. *J Pediatrics* 1991; 119:417-423.

<http://ballardscore.com/Pages/ScoreSheet.aspx>